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Natural products influence bacteriophage infectivity

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Bacteriophages (phages) are obligate viruses that infect bacteria. The antibacterial effects of both phages and natural products shape microbial ecosystems and have yielded competing antibiotic strategies. Phages have also intersected many times with natural products research throughout the past century. To discover antiviral leads, natural products were screened for anti-phage activity. To discover new anti-cancer drugs, natural products were screened for the ability to trigger lysis by the λ prophage—indicating DNA damage. Now, the antibiotic resistance crisis motivates the study of natural products that can synergize with phages to improve antibacterial therapies. Beyond applications, these parallel natural “chemical” and “biological” antibacterial factors combine to shape microbial communities across our planet. Here, we provide a comprehensive overview of natural products that modulate phage activities. We discuss their mechanisms of action, and we present opportunities for future research.

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7. **Conflicts of interest**
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1. Introduction

Frederick Twort and Félix d'Hérelle discovered bacteriophages (phages) over a century ago.¹ Since then, the study of phages has been pivotal for the fields of virology, molecular biology, microbial ecology, and antibacterial therapeutics. As viruses that exclusively infect (and often lyse) bacteria, phages are natural killers of pathogenic bacteria. Shortly after Félix d'Hérelle identified phages, he realized their therapeutic potential and explored the possibility of using phages to treat bacterial infections in both animals and humans.² His early efforts in “phage therapy” pioneered the way bacterial infections are treated today in some parts of the globe—perhaps most notably in the nation of Georgia.³ Phage research in the 20th century also led to paradigm-shifting discoveries⁴ including the realization that DNA is nature's hereditary material,⁵ the identification of mRNA as the short-lived intermediate before protein synthesis,⁶ the employment of restriction enzymes in molecular biology,⁷ the development of phage display techniques to identify countless peptide-binding interactions,⁸ and the employment of CRISPR-Cas in genetic engineering.⁹ Furthermore, the recent antibiotic resistance crisis¹⁰ is reviving global interest in phage therapy.¹¹ This renewed excitement in phage research warrants a review of the long history of natural products' influence on phages, as well as a discussion of recent discoveries and avenues for future research.

The two major life cycles found in phages are the lytic cycle and the lysogenic cycle (Fig. 1). Obligately lytic phages only undergo the lytic cycle, kill the host, and release new progeny to the environment (Fig. 1). In contrast, temperate phages can

undergo both the lytic and the lysogenic cycles. In the lysogenic cycle, phages integrate their DNA into the bacterial genome and lay dormant within the host as a prophage. When the right conditions arise, the prophage can excise from the host genome and undergo the lytic pathway to infect nearby cells (Fig. 1).

Studies on natural products that influence phage behavior have led to the discovery of anti-viral compounds,¹² anti-cancer drugs,^{13–15} and phage-antibiotic synergies.^{16–19} Beyond these medical applications, secondary metabolites also shape the natural symbiotic relationships between microbial species by modulating phage activities. On one hand, anti-phage molecules produced by one species may protect a polymicrobial community from phage predation, thus fostering a mutualistic or commensal interaction. On the other hand, metabolites may promote phage infections or induce lysogenic phages into the lytic cycle, which could benefit the metabolite producer by eliminating its competing bacteria.

A revived interest in phages is refocusing some chemists on the interactions of natural products with phages. Phage-promoting natural products may be co-administered with phages to improve phage therapy. In contrast, natural products in the environment or host may inhibit phage therapy, and therefore necessitate alternate strategies. Finally, phage-metabolite synergies may shape microbiome health. These applications justify a deep exploration into the known interactions between natural products and phages (and call for further research to expand the current frontiers).

We note that other recent reviews have discussed small molecules that inhibit phage infections and affect lysis-lysogeny decisions.^{20–22} This review goes beyond to also include small molecules that promote phage infections. As our topic focuses on natural products, synthetic or semisynthetic compounds will be largely excluded from discussion.

This review is organized primarily by the categories of impact on phages. First, we discuss natural products that inhibit phage proliferation. Then, we discuss metabolites that promote phage



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Zhiyu Zang received his BSc degree in chemistry at Peking University, China. He then moved to the United States and obtained his PhD degree at Indiana University Bloomington under the supervision of Prof. Joseph P. Gerdt. His doctoral research focused on the intersection of chemistry and microbiology, specifically exploring the modulation of phage infection and anti-phage systems by natural products and synthetic small molecules.



Joseph P. Gerdt

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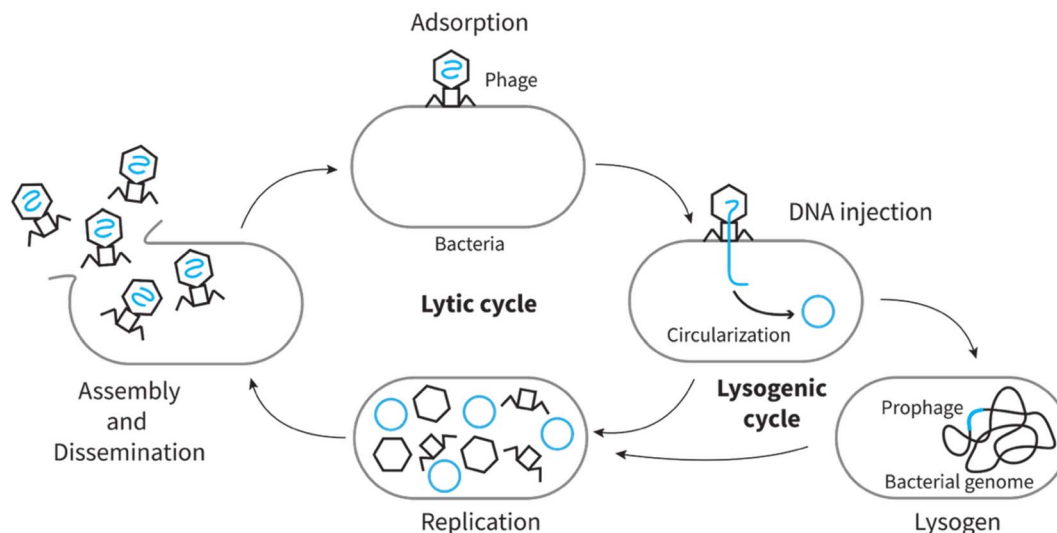


Fig. 1 The lytic and lysogenic life cycle of phage.

replication on bacterial hosts. Finally, we discuss natural products that impact the lysis–lysogeny decision of temperate phages. Each category is further divided by the mechanisms by which the natural products carry out their influences on phages. In some cases, the mechanisms are still poorly understood, but we do our best to explain the likely modes of action, given insights from non-phage studies.

2. Anti-phage natural products

Dozens of phage-inhibiting natural products have been described over the decades. These discoveries have implications for anti-viral therapy and microbial ecology. Since a molecule that inhibits bacteriophage replication sometimes also inhibits viruses that infect animals,^{12,23,24} anti-phage natural products provide an easy initial screen for the discovery of new anti-viral compounds. Furthermore, in nature, these anti-phage natural products likely shape microbial ecosystems. The evolutionary pressures driving the production of anti-phage natural products are debatable. One hypothesis is that microbial-encoded anti-phage natural products might have evolved as immune mechanisms against phage attacks.^{25,26} Beyond self-immunity, these anti-phage metabolites might also provide “herd immunity” against phage predation for an entire microbial community. However, as discussed below, many (but not all) anti-phage natural products are also antimicrobial. Therefore, it is possible that the production of several anti-phage natural products was primarily driven by their direct influence on microbial competitors—not their anti-phage activity. Regardless of their evolution, anti-phage metabolites have the capacity to shape microbial ecology. They may also diminish the efficacy of phage therapy, warranting attention to the complex chemical environments that can influence phage–bacteria interactions.

Multiple methods have been employed to assess the anti-phage activity of natural products. We highlight two methods (Fig. 2) that can reveal selective anti-phage activity by molecules

that are not antimicrobial (at least at the applied dose). One case monitors the reduction of plaques (areas of phage-induced bacterial lysis on an agar surface). The other case monitors a reduction of phage-induced lysis in liquid culture.

Most anti-phage natural products arrest core phage functions that are also core cellular functions, such as DNA replication, transcription, and protein synthesis. In many cases, anti-phage molecules inhibit phage proliferation more potently than host cell replication. This selectivity may stem from phage-specific molecular targets (*e.g.*, linear DNA and phage-encoded enzymes) being more sensitive to anti-phage molecules. Alternatively, because phages are fast-replicating entities, they may simply be more susceptible to minor perturbations. Apart from inhibiting core functions of genome replication and gene expression, some anti-phage natural products inhibit phage attachment by inducing modifications to the host cell surface. Below, we discuss individual anti-phage natural products, categorized by their likely mechanisms of action.

2.1. Interfere with DNA replication and/or transcription

DNA-binding molecules are the most commonly observed anti-phage natural products. These molecules antagonize phage reproduction by interfering with phage DNA synthesis and/or transcription (Fig. 3). The binding of these molecules to DNA may stall the movement of DNA/RNA polymerase along DNA or inhibit the coiling and relaxing of DNA by topoisomerases (Fig. 3).²⁷ Following are examples of anti-phage natural products that bind DNA.

2.1.1. Anthracyclines. Anthracyclines make up a class of antibiotics and chemotherapy drugs. They contain a tetracyclic backbone with an anthraquinone core and a sugar moiety (Fig. 4A). This class of molecules has an extensive research record—primarily due to the antitumor activities of many of its members. The first molecule in this class with reported anti-phage activity was aklavin in 1955, which was purified from *Actinomyces*.²⁸ It was shown to inhibit phages T2 and T5



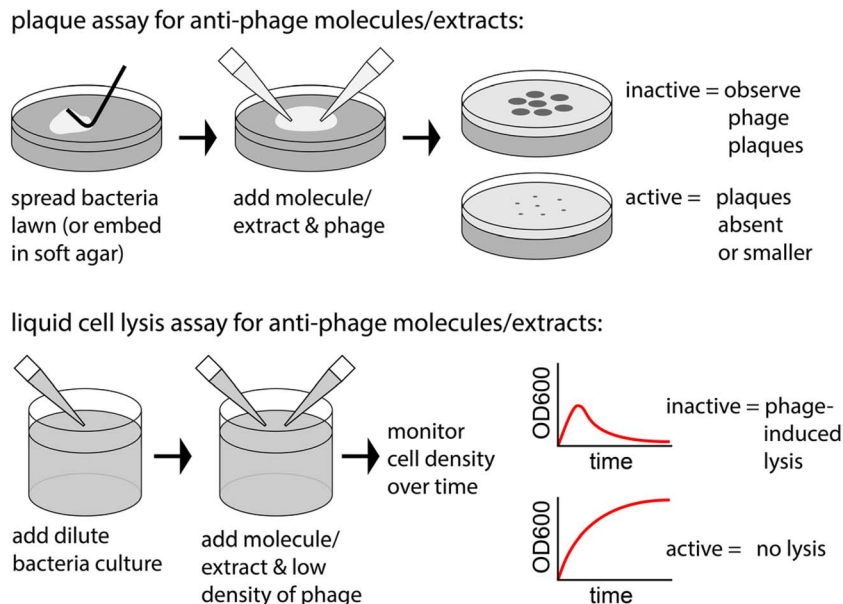


Fig. 2 Example experiments to observe anti-phage natural products.

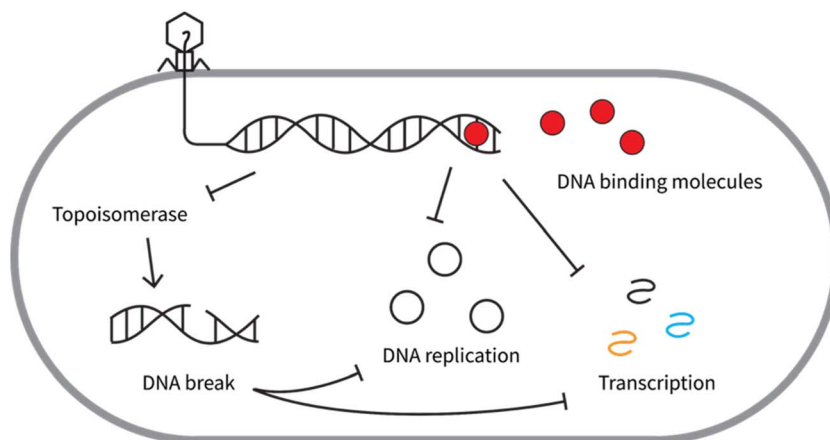


Fig. 3 The major mechanisms of action of anti-phage DNA-binding molecules.

forming plaques on *Escherichia coli*, as well as a diverse panel of phages infecting other bacteria.²⁸ Later studies in the 1960s and 1970s reported that other molecules belonging to the anthracycline family could specifically inhibit DNA phages but not RNA phages. These selective inhibitors include daunorubicin²⁹ (*i.e.*, daunomycin, isolated from *Streptomyces peucetius*³⁰), doxorubicin³¹ (*i.e.*, adriamycin, isolated from *Streptomyces peucetius*³²), and aclarubicin³¹ (*i.e.*, aclacinomycin A, isolated from *Streptomyces galilaeus*³³).

The specific inhibition of DNA phages suggested a favorable interaction between anthracyclines and DNA. Indeed, anthracyclines interact with DNA by intercalating their planar tetracycline backbone between two alternating C–G base pairs with the amino sugar extended into the minor groove of the DNA double helix, as visualized by X-ray diffraction (Fig. 4B).^{34–37} Because anthracycline binding can stabilize the DNA duplex, these molecules may inhibit phage infection by directly

interfering with the action of both DNA polymerase³⁸ and RNA polymerase³⁹ (Fig. 3).

Another possible mechanism to explain the inhibition of anthracyclines on phage DNA replication and transcription is that they can interfere with the function of type II topoisomerases (Fig. 3).⁴⁰ Type II topoisomerase is an important enzyme for DNA replication and transcription during phage infection. It cuts both strands of the DNA helix and reseals them to manage DNA tangles and supercoils.⁴¹ For example, T-even phages encode their own type II topoisomerase, which is required to relax potential DNA supercoils or to resolve DNA knots of the rapidly replicating genome.^{42,43} With anthracycline intercalated into DNA, a stable ternary complex forms among anthracycline, DNA, and topoisomerase, which prevents the ligation of double-stranded DNA breaks (DSBs) by the topoisomerase.⁴⁴ Anthracyclines might also induce DSBs through a radical mechanism due to the presence of quinone moiety.⁴⁵ These irreversible DSBs



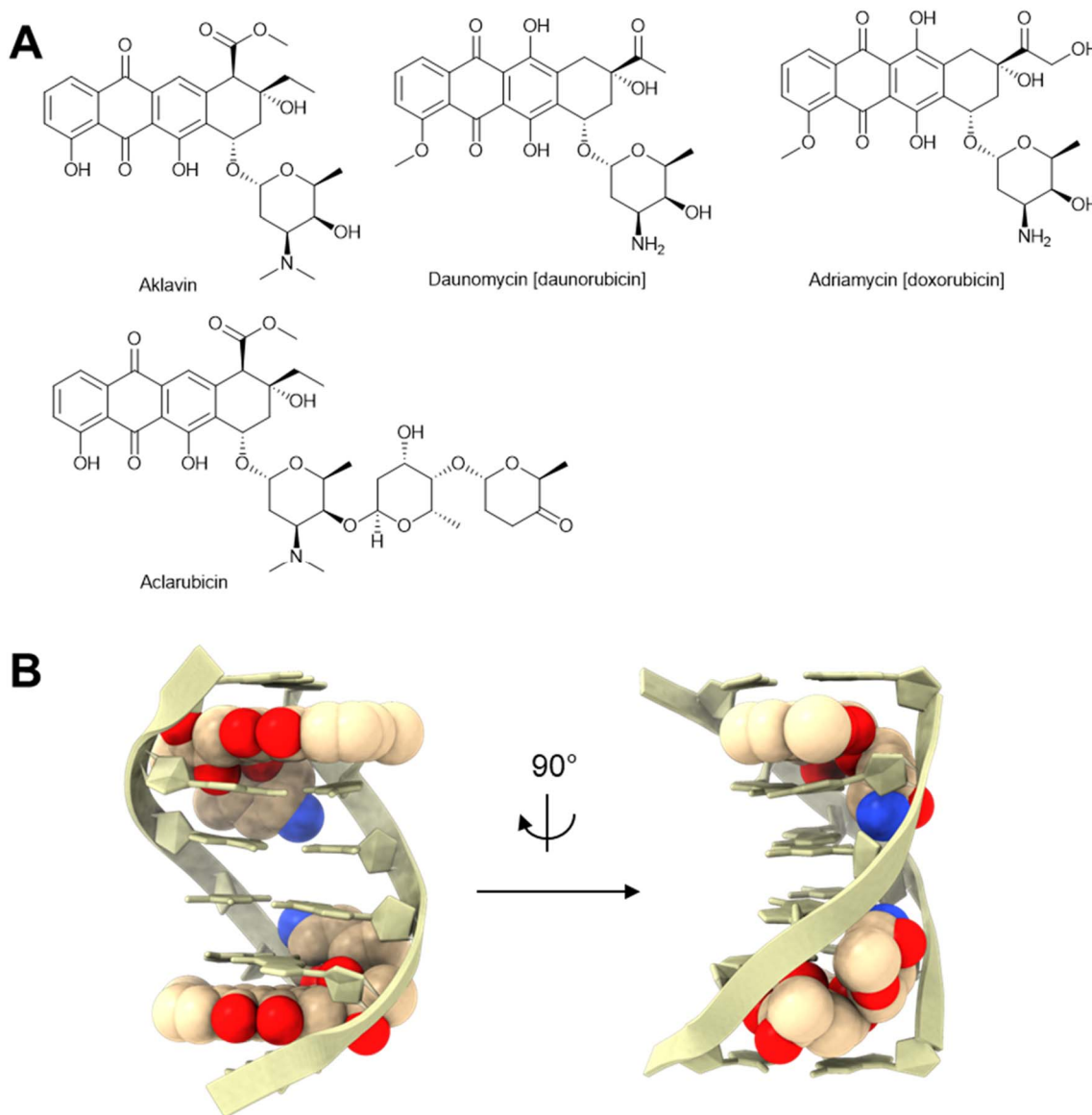


Fig. 4 Anthracycline anti-phage molecules. (A) Chemical structures of anthracyclines. (B) Daunorubicin:DNA complex [PDB: 1D10].

inhibit DNA replication and transcription (Fig. 3),⁴⁶ presumably hampering the phage infection process.

With several potential mechanisms, there is still uncertainty about how anthracyclines inhibit phage infection. As for lysogenic phages, a recent study showed that anthracyclines did not prevent the λ phage genome from entering the cell but significantly reduced the ability of the phage genome to integrate into the bacterial chromosome as a lysogen.²⁵ The exact mechanism by which anthracyclines inhibit lysogen formation is still unclear. Surprisingly, anthracyclines did not inhibit λ phage replication after induction of the temperature-sensitive λ prophage.²⁵ This discovery complicates the phage inhibition mechanism of anthracyclines, because they do not seem to universally inhibit DNA replication and transcription of all phages. Future investigations on the interaction between anthracyclines and phages may unravel the mechanism behind their selective anti-phage activity.

Although anthracyclines are also anti-bacterial, the phage genome is suspected to be more susceptible to DNA intercalators compared to the bacterial genome, partly because phage DNA is linear, non-supercoiled, and unprotected by DNA-binding proteins when it is injected into the bacterial host.²⁵ Therefore, at low doses, anthracyclines can selectively inhibit phages more than their host bacteria.^{25,29,31}

2.1.2. Neopluramycin. Neopluramycin was first isolated from *Streptomyces pluricolorescens* in 1970 and exhibited antibiotic and anticancer activities.⁴⁷ Shortly after its discovery, neopluramycin was also found to inhibit the production of T4 phage particles in *E. coli*.⁴⁸ The phage inhibition activity was due to interference with phage transcription as measured both *in vitro* and in infected cells.⁴⁸ Neopluramycin has a tetracyclic backbone similar to anthracyclines but with a pyran ring fused to the anthraquinone chromophore (Fig. 5). The planar backbone presumably allows neopluramycin to intercalate between



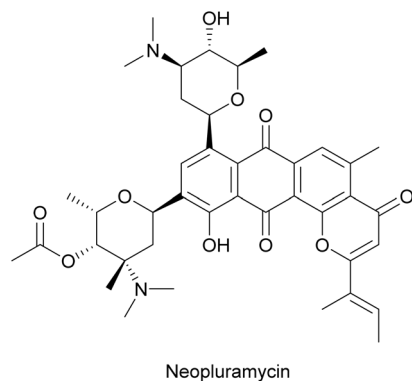


Fig. 5 Chemical structure of neopluramycin.

two adjacent base pairs with the two amino sugars residing in the minor groove as inferred by NMR studies on its analogue, hedamycin.^{49–51} As with the anthracyclines, the antiphage effect of neopluramycin probably results from its DNA-intercalating properties, which may not only interfere with phage transcription but also with DNA synthesis (Fig. 3).

2.1.3. Gilvocarcin family of C-aryl glycosides. The gilvocarcin C-aryl glycosides (also referred to as benzo[*d*]naphtho[1,2-*b*]pyran-6-one C-glycosides) are known for their excellent antitumor activity and remarkably low toxicity.⁵² This family of natural products contains a tetracyclic naphthocoumarin backbone and a vinyl substituent at the C8 position, with various sugars attached to the C4 position of the aromatic backbone *via* a C–C bond

(Fig. 6A). Chrysomycin A was the first molecule discovered within this family. It was isolated from a *Streptomyces* bacterium in 1954.⁵³ Chrysomycin A inhibits plaque formation by a variety of phages, including coliphages T1 and T2, *Bacillus* phages, *Staphylococcus* phages, and *Enterococcus* phages.⁵³ Another member in this family, gilvocarcin V (toromycin), was also shown to inhibit phage infection by phi170, T1, T3, and T5 phages in 1979. The same study reported inhibition of DNA viruses of animals like the vaccinia virus and the herpes simplex virus, but not RNA viruses like the Newcastle disease virus.²³ Later it was shown that gilvocarcin V can bind to single-stranded DNA of coliphage M13 *in vitro*.⁵⁴

The mechanism of inhibition of the gilvocarcin family against phages is still unclear. We speculate that they hinder phage replication by inhibiting phage DNA synthesis through photo-activated DNA alkylation (Fig. 6B). The alkylation relies on visible light or low energy UV radiation, which initiate a [2 + 2] photocycloaddition between the gilvocarcin vinyl group and thymine residues of DNA (Fig. 6B),⁵⁵ thereby inhibiting DNA synthesis and causing DNA damage.^{56,57} Moreover, gilvocarcins might interfere with phage DNA synthesis and transcription through the inhibition of topoisomerases (Fig. 3), as chrysomycin A has been shown to inhibit the activity of both type I and type II topoisomerase, presumably through binding to DNA and/or blocking the topoisomerase active sites.^{58,59}

Beyond the phage inhibition activity of gilvocarcin family molecules, they can also trigger prophage induction by causing extensive DNA damage in host cells,⁶⁰ which is discussed later in the prophage induction section.

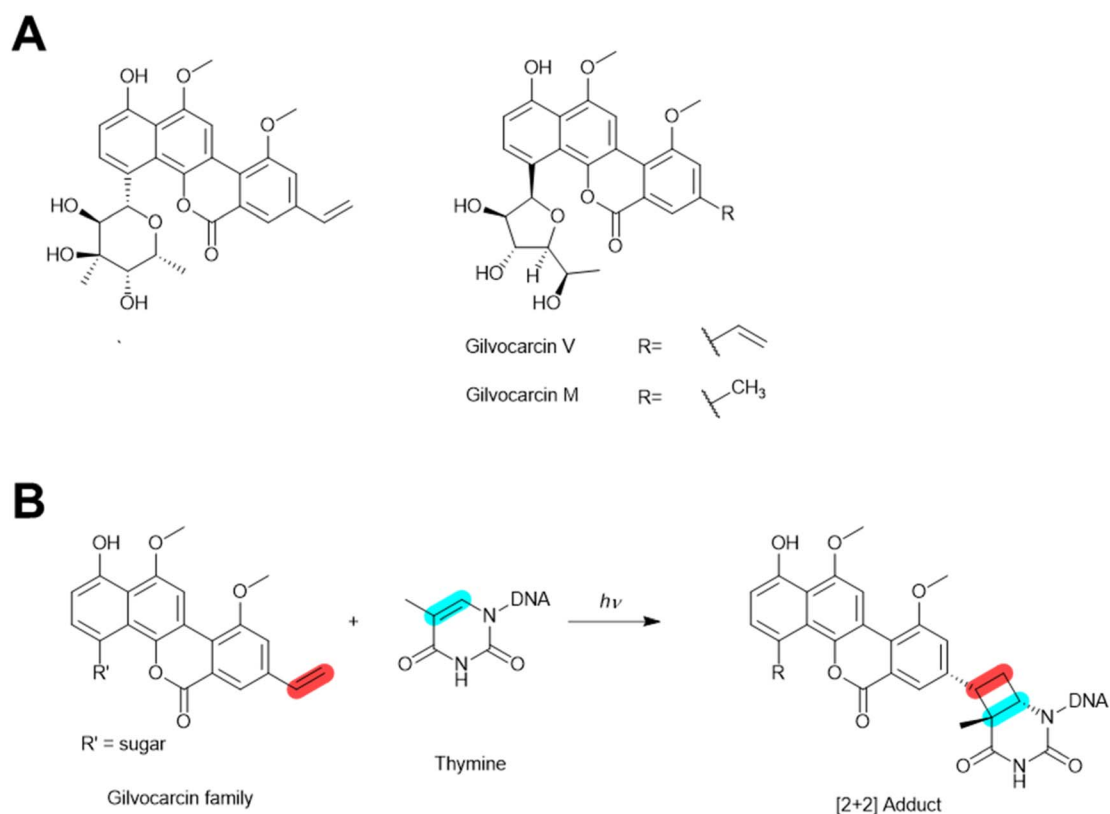


Fig. 6 Gilvocarcin anti-phage molecules. (A) Chemical structures of gilvocarcins. (B) Photo-activated DNA alkylation by gilvocarcins.



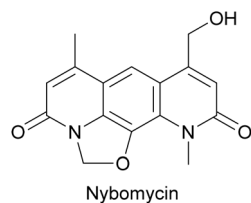


Fig. 7 Chemical structure of nybomycin.

2.1.4. Nybomycin. Nybomycin is a pyridoquinolinedione-based antibiotic first collected from an Actinomycetia isolate in 1955 (Fig. 7).⁶¹ The authors tested its ability to restrict plaque formation by a panel of phages and found 33 out of 61 phages were inhibited by nybomycin, including phages of *E. coli*, *Staphylococcus*, *Bacillus*, and *Streptomyces*.⁶¹

Although the exact mechanism of phage inhibition by nybomycin is still unclear, it likely acts *via* DNA intercalation and/or inhibition of type II topoisomerase. Molecular docking shows that its planar pyridoquinolone structure enables nybomycin to partially intercalate into a DNA double helix.⁶² Additionally, nybomycin can inhibit type II topoisomerase *in vitro* (presumably by stabilizing the nicked DNA-topoisomerase complex), which disrupts DNA supercoiling and relaxation.⁶² Therefore, it is likely that nybomycin's DNA intercalation and/or topoisomerase inhibition prevents phage DNA replication and/or transcription as discussed above (Fig. 3).

2.1.5. Oligopyrrole/polyamide. Netropsin (*i.e.*, T-1384, congocidine, or sinanomycin) and distamycin A are two naturally occurring amide-linked oligopyrrole antibiotics (Fig. 8A) isolated from *Streptomyces netropsis*⁶³ and *Streptomyces distallicus*,⁶⁴ respectively in the 1950s. Distamycin A was reported to inhibit phage T1 (ref. 65) and T2 (ref. 66) infection in *E. coli* as

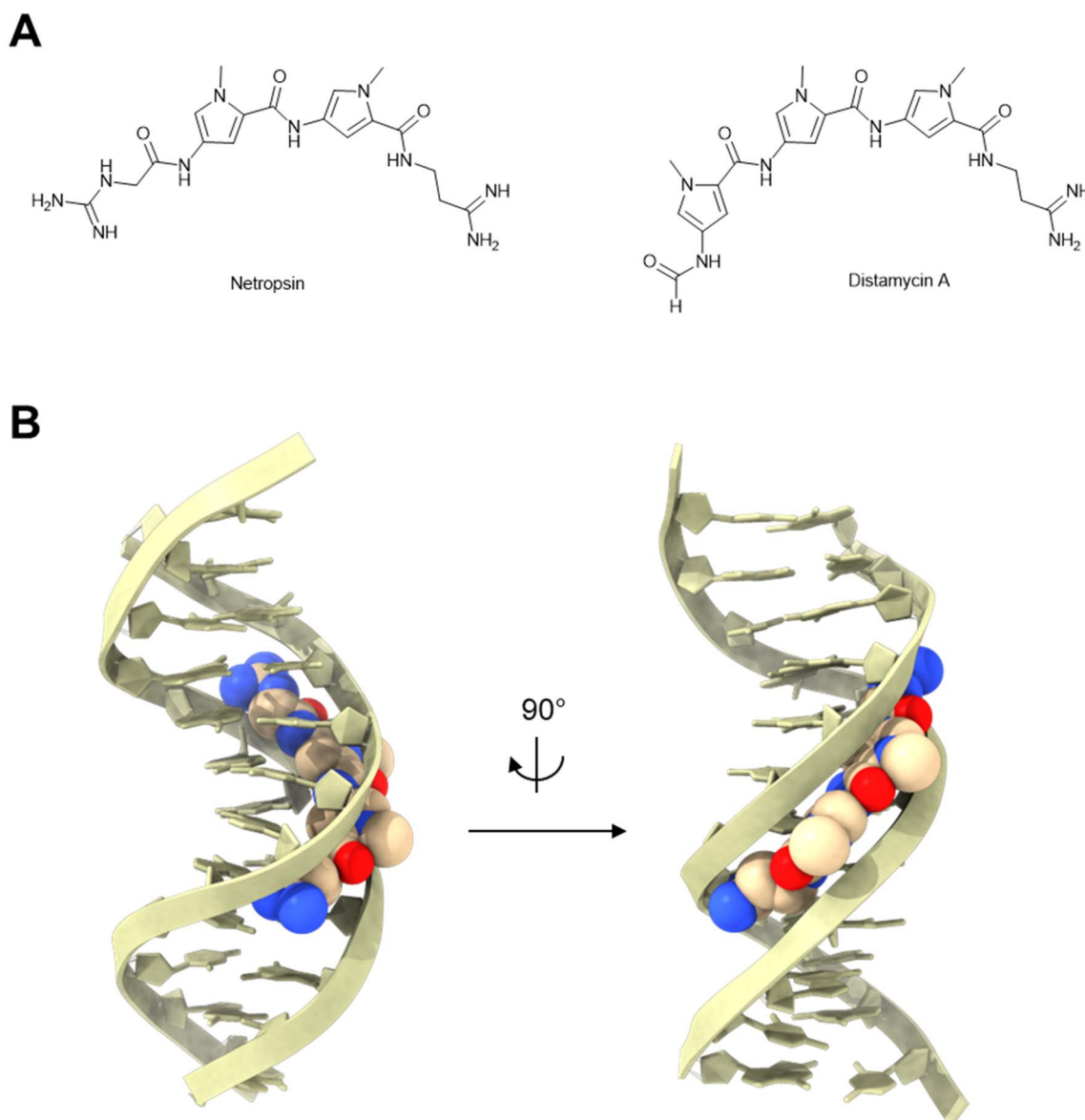


Fig. 8 Oligopyrrole anti-phage molecules. (A) Chemical structures of oligopyrroles. (B) Netropsin:DNA complex [PDB: 6BNA].



evidenced by impeded plaque formation and protection from phage-induced host culture lysis.⁶⁵ Although distamycin A can arrest bacterial growth, it selectively inhibited phage replication at low concentrations that do not inhibit growth of the host bacteria.⁶⁵ Notably, the anti-phage activity of distamycin A also inspired its subsequent investigation as an inhibitor of animal viruses.¹² Netropsin has not been tested against phages. However, it inhibits the proliferation of several animal viruses, such as vaccinia,⁶⁷ influenza,⁶⁸ and Shope fibroma.⁶⁹ Therefore, it is likely to inhibit phages, as well.

The anti-phage effect of oligopyrrole antibiotics is presumably due to their specific binding within the minor groove of the DNA double helix (Fig. 8B).^{70–72} The binding of oligopyrroles to DNA can interfere with phage DNA replication and transcription in multiple ways. First, distamycin A has been shown to directly inhibit both DNA⁷³ and RNA⁷⁴ synthesis *in vitro*.⁷⁵ Phage transcription is preferentially inhibited by distamycin A, compared to bacterial transcription. Namely, a direct *in vitro* comparison revealed that distamycin A inhibited phage T3 RNA polymerase more strongly than an *E. coli* RNA polymerase.⁷⁶

This discovery is in agreement with the selectivity of distamycin A to inhibit phage replication at concentrations that do not arrest bacterial growth.⁶⁵ Second, the binding of distamycin to the DNA minor groove can prevent the catalytic activity of both type I⁷⁷ and type II⁷⁸ topoisomerases by blocking the enzyme binding sites. As discussed earlier, topoisomerase inhibition can indirectly interfere with DNA replication and transcription, thus blocking rapid phage replication (Fig. 3).

2.1.6. Actinomycins. Actinomycins are a class of chromopeptides with potent cytotoxicity and antimicrobial activity.⁷⁹ Actinomycins feature a phenoxazinone chromophore tethered to two cyclic pentadepsipeptides *via* amide bonds (Fig. 9A). It was first reported in 1961 that a mixture of actinomycins (referred to as actinomycin S) isolated from *Streptomyces flavovolus* 1048A⁸⁰ inhibited multiplication of phage T2 on *E. coli* while not interfering with host growth.⁸¹ A following study showed that the two major components in actinomycin S were actinomycin D and actinomycin X₂ (Fig. 9A).⁸² Later, it was shown that actinomycin D can inhibit infections by other phages as well, including *E. coli* phage T4,⁸³ and *Bacillus subtilis*

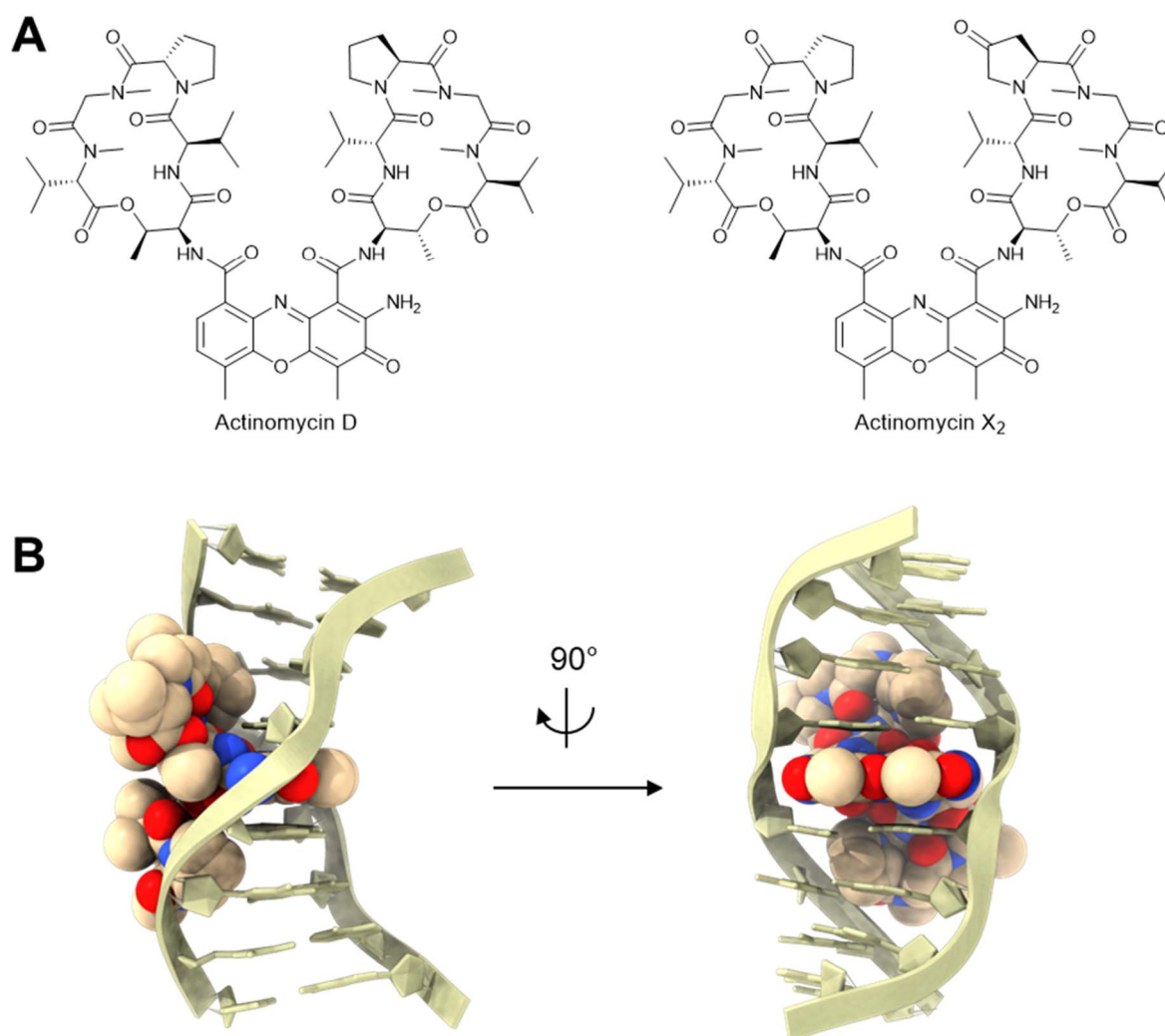


Fig. 9 Actinomycin anti-phage molecules. (A) Chemical structures of actinomycins. (B) Actinomycin D:DNA complex [PDB: 2D55].



phages PBS1 and SP10.⁸⁴ Around the same time, actinomycin D was also shown to inhibit the single-stranded RNA virus that causes foot-and-mouth disease in animals.⁸⁵

The planar structure of tricyclic phenoxazinone in actinomycins allows them to intercalate selectively between alternating G–C base pairs, while the two cyclic pentadepsipeptides

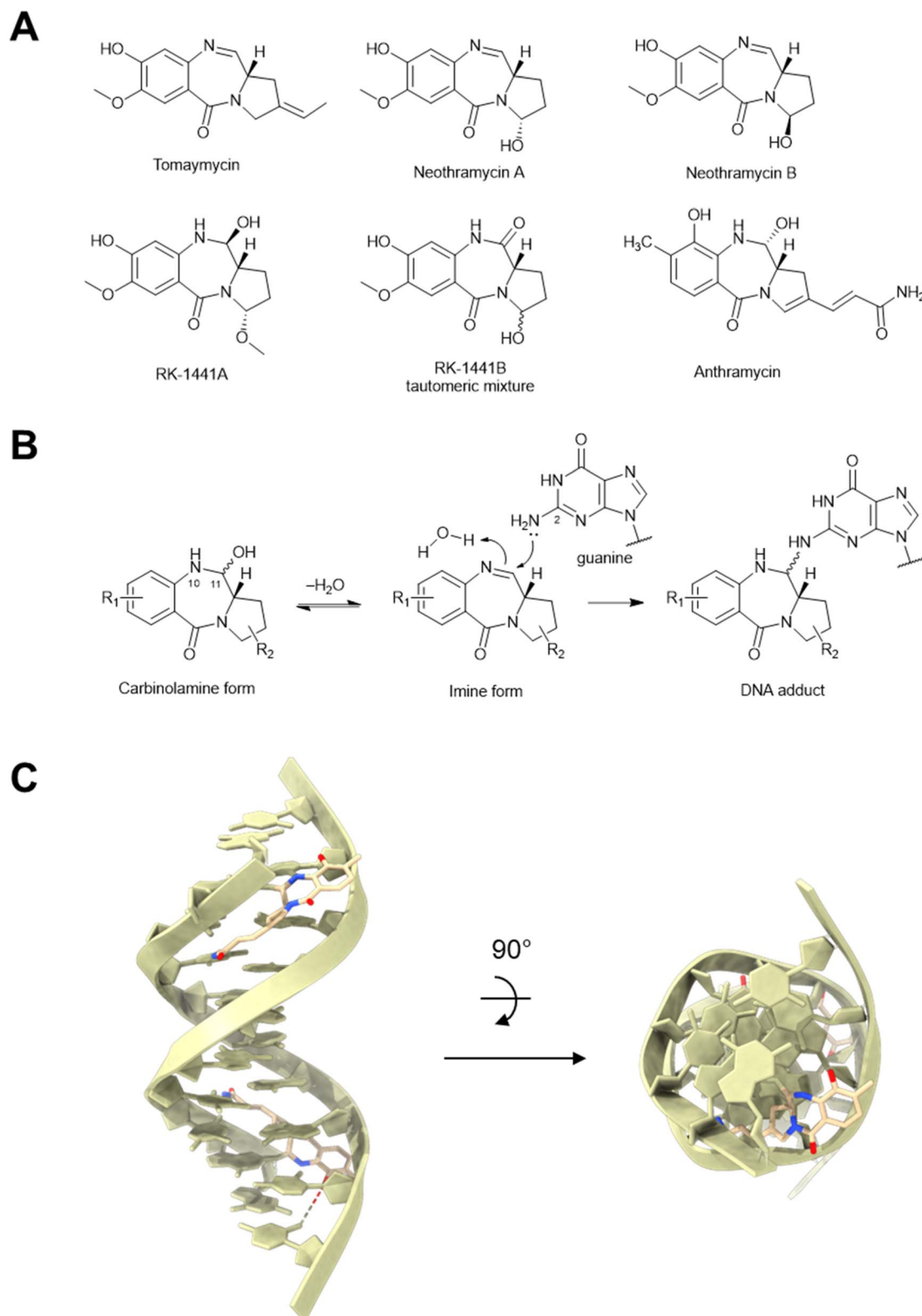


Fig. 10 Anthramycin anti-phage molecules. (A) Chemical structures of anthramycins. (B) Anthramycin DNA alkylation mechanism (C) Anthramycin-DNA adduct [PDB: 274D].



imine intermediate and then alkylate DNAs. The S configuration at C-11a confers pyrrolobenzodiazepines a right-handed twist from the anthranilate to the hydroxyproline ring, allowing them to fit perfectly in the minor groove of the DNA double helix (Fig. 10C).¹⁰⁰ It is surprising that RK-1441B was also active against phage infections, as the stable amide between N-10 and C-11 is not reactive *in vitro* with purified phage DNAs.⁹⁶ It was proposed that the amide might be converted into the active carbinolamine or imine form in the host cells,⁹⁶ which would allow RK-1441B to alkylate DNA as well.

2.1.8. Lanthipeptide. Lanthipeptides are one of the largest and most diverse families of ribosomally synthesized and post-translationally modified peptides (RiPPs).¹⁰² Lanthipeptides are named after the presence of their characteristic β -thioether linked bis-amino acid structures, lanthionine (Lan) and methylanthionine (MeLan) (Fig. 11A), which are formed by cysteine residues crosslinking with dehydrated serine or threonine residues, respectively.¹⁰³ Lanthipeptides are notable for their antimicrobial,¹⁰⁴ anti-cancer,¹⁰⁵ and anti-animal virus¹⁰⁵ properties. Recently the first anti-phage lanthipeptide was reported.²⁶ Through bioinformatics analysis of the genomes from Actinomycetota, the authors discovered that lanthipeptide biosynthetic gene clusters (BGCs) reside near other anti-phage systems within defense islands (Fig. 11B) at a frequency of 8.8%.²⁶ This observation suggested that these lanthipeptides serve as anti-phage defenses for the host.^{106,107} Indeed, upon induced native expression or heterologous expression, lanthipeptide BGCs provided robust protection against phage infections in *Streptomyces* (Fig. 11C, predicted core peptides are shown, intramolecular β -thioether linkages are yet uncharacterized).²⁶ The lanthipeptide inhibited phage transcription, particularly the late genes.²⁶ By comparing the genomes of wild type phages with their lanthipeptide-immune escape mutants, the authors discovered that each of the escaping phages carried a mutant Gcn5-related *N*-acetyltransferase (GNAT). Phage-encoded GNATs are important for shifting between early and late gene expression.¹⁰⁸ Therefore, the lanthipeptide might inhibit transcription through a yet unknown GNAT-dependent mechanism (Fig. 11D).²⁶ It is noteworthy that only

intracellular lanthipeptides were found active against phage infections in *Streptomyces* so far.²⁶ Therefore, further investigation is required to determine if secreted lanthipeptides from one bacteria can inhibit phage infection in another bacteria. If not, these peptides may only be retained within the producing cell for its own defense.

Perhaps more than any other anti-phage natural product, the main purpose of these lanthipeptides appears to be anti-phage defense. Their biosynthetic genes are located in “phage defense islands”, and the lanthipeptides do not exhibit obvious antibiotic activity. This case contrasts with the previously discussed anti-phage natural products that are also antibiotics. The dual anti-phage/antibiotic activity of the other molecules adds to an existing debate about the evolved roles of naturally occurring antibiotics to benefit the producing organism. As others have noted, antibiotics may mediate microbial competition by killing competitors, or they may serve as signal molecules to regulate transcriptional profiles.¹⁰⁹ Here we note the third possibility: antibiotics with anti-phage activities might have evolved as immune mechanisms against phage infections. Despite this debate, the case of anti-phage intracellular lanthipeptides appears fairly clear—they likely evolved for defense against phages.

2.2. Interfere with peptide synthesis

Peptide synthesis inhibitors comprise another large group of anti-phage natural products. These molecules target the bacterial ribosome. Since phage protein synthesis exclusively relies on host ribosomes,¹¹⁰ the inhibitors of host ribosomes also interfere with the synthesis of phage-encoded peptides, thereby reducing phage reproduction (Fig. 12).

2.2.1. Aminoglycosides. Aminoglycosides were among the first antibiotics to be introduced for clinical treatment of bacterial infections, and they remain one of the major classes of antibiotics in use today.^{111,112} Aminoglycosides feature a set of sugars, amino sugars, and pseudo sugars (*e.g.*, cyclitols and aminocyclitols) that are connected *via* glycosidic linkages (Fig. 13). Streptomycin was the first reported aminoglycoside, isolated from *S. griseus* by Selman Abraham Waksman and

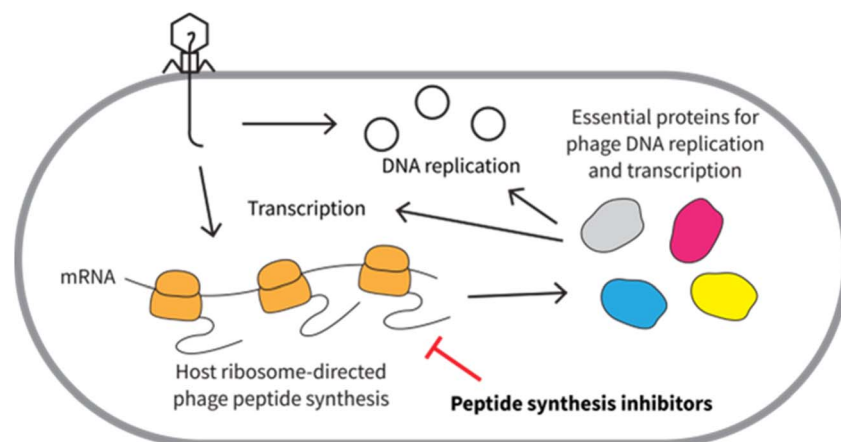


Fig. 12 Hypothetical anti-phage mechanisms of peptide synthesis inhibitors.



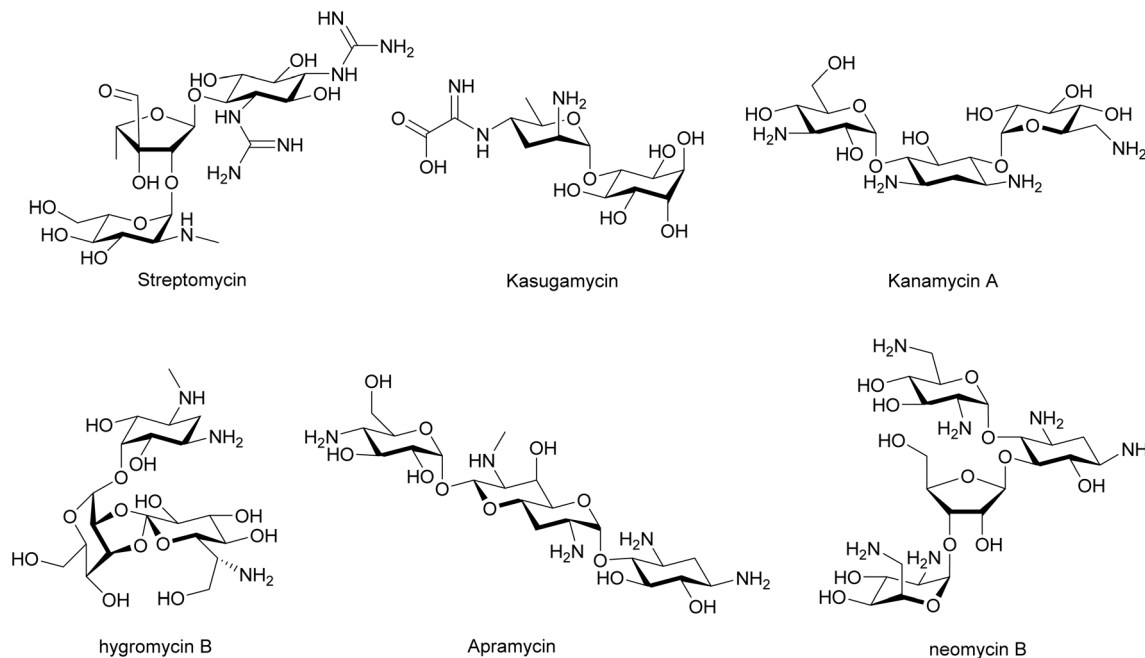


Fig. 13 Chemical structures of anti-phage aminoglycosides.

colleagues in 1944.¹¹³ Soon after its discovery, streptomycin was reported to suppress plaque formation and phage multiplication in both *E. coli* and *Staphylococcus aureus* in 1945.¹¹⁴ In the following decades, the anti-phage effect of streptomycin was

demonstrated against many other phages.^{115–120} Other aminoglycosides beyond streptomycin, such as kasugamycin,¹²¹ kanamycin A,^{120,122,123} hygromycin B,^{120,123} apramycin,¹²³ and

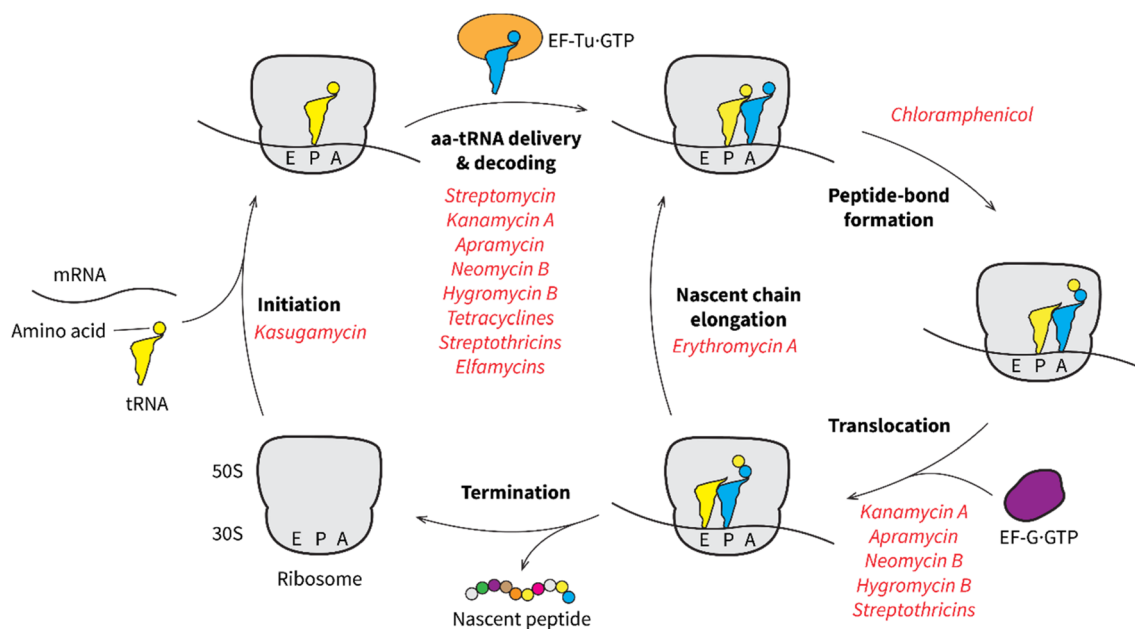


Fig. 14 Scheme of peptide synthesis and ribosome recycling. Peptide synthesis is initiated by the formation of a complex between the 70S ribosome (including a small 30S subunit and a large 50S subunit), mRNA, and the initiator tRNA at the P-site. The elongation cycle involves four steps. First, an aminoacylated tRNA (aa-tRNA) is delivered to the A-site with the help of elongation factor Tu (EF-Tu). Upon delivery, the ribosome ensures the correct pairing between the tRNA anti-codon and the mRNA codon (decoding). Next, the amino acid (or peptide in the elongation process) from the P-site tRNA is transferred to the aa-tRNA at the A-site, and a peptide bond is formed. The ribosome-tRNA complex then translocates to the next codon on the mRNA with the help of elongation factor G (EF-G), so that the next aa-tRNA can be delivered to the A-site. In the elongation process, the nascent peptide chain passes through the exit tunnel in the 50S subunit. The elongation cycle terminates when a stop codon is encountered and the nascent peptide chain is released from the ribosome. Steps that are inhibited by natural products are indicated.



neomycin B,^{122,124} have also proven active against phage infection.

The mechanism of antiphage activity from aminoglycosides can be related to their specific interaction with the 30S or 50S subunits of the bacterial ribosome, thus inhibiting distinct steps in protein translation, such as tRNA delivery and selection, ribosome translocation, and ribosome recycling (Fig. 14).^{125–132} As host ribosomes are essential for phage protein synthesis,¹¹⁰ malfunction of host ribosomes should interfere with production of phage proteins. It is plausible that the higher demands of phages for fast replication makes them even more susceptible than their hosts to subtle ribosome inhibition by low concentrations of aminoglycosides, affording anti-phage functions at sub-inhibitory doses.

A recent study proposed another anti-phage mechanism of aminoglycosides by directly inhibiting phage DNA replication and transcription.¹²³ The authors discovered that apramycin treatment led to a significant reduction in phage DNA replication and transcription.¹²³ Since *in vitro* studies have shown that aminoglycosides can cause condensation of purified phage DNA,¹³³ the authors suggested that the impaired phage DNA replication and transcription was due to direct binding of aminoglycosides to the unprotected phage DNA following the injection.¹²³ Alternatively, the decreased phage DNA and RNA synthesis might result from the impaired function of host ribosomes under aminoglycoside treatment. Phage-encoded proteins are often vital for efficient phage DNA and RNA synthesis. They arrest host gene expression, redirect host DNA and RNA polymerases to phage genomes, assist the initiation of DNA replication, and regulate transcription kinetics.^{134–138} Since phages rely on the host ribosomes for their protein synthesis,¹¹⁰ the inhibitory actions of aminoglycosides on the host ribosomes may be the root cause of the observed decrease in phage DNA replication and transcription (Fig. 12).

2.2.2. Tetracyclines. Tetracyclines are a class of broad-spectrum antibiotics characterized by a rigid fused tetracyclic core with a variety of functional groups attached.¹³⁹ In 1948, the first molecule in this class was isolated from *Streptomyces aureofaciens*, named aureomycin (*i.e.* chlortetracycline, Fig. 15).¹⁴⁰ A few years later, chlortetracycline was shown to inhibit phage T3 infection in *E. coli* by slowing down phage reproduction and reducing its burst size (the number of new phages produced by each infected cell).²⁴ In addition, the authors showed that chlortetracycline inhibited phage adsorption onto the host bacteria.²⁴ As with many other anti-phage metabolites, chlortetracycline also inhibited an animal virus.¹⁴¹ The non-

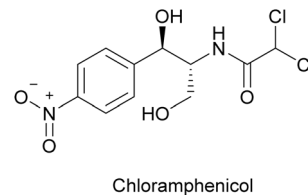


Fig. 16 Chemical structure of chloramphenicol.

chlorinated analog, tetracycline (Fig. 15), was also reported to inhibit the T3 phage recently.¹²² In that report, tetracycline did not inhibit phage adsorption. The different effects of tetracyclines on phage adsorption is intriguing because chlortetracycline and tetracycline only differ by a chloro group. Further investigation may be warranted to elucidate the importance of the chloro group in antagonizing phage adsorption. Nonetheless, the consistent inhibitory effect of tetracycline on phage reproduction is probably due to the impaired ribosomal function as tRNA delivery is inhibited.^{142,143}

The anti-phage effect of tetracycline also intersects with a bacterial defense system (CRISPR-Cas) and a phage-encoded anti-CRISPR (Acr) system. In one case, bacteriostatic antibiotics like tetracycline, chloramphenicol, and erythromycin, promoted CRISPR immunity in a *P. aeruginosa* population by slowing the phage maturation process, thus allowing more time for spacer acquisition.¹⁴⁴ In another case, the infection of CRISPR-containing *P. aeruginosa* by Acr-encoding phages was also inhibited by tetracycline and the other translation inhibitors chloramphenicol and erythromycin.¹⁴⁵ These translation inhibitors delayed the production of phage-encoded “immunosuppressing” Acr proteins, thereby allowing the CRISPR immune system to inhibit phage reproduction.¹⁴⁶

2.2.3. Chloramphenicol. Chloramphenicol (chloromycetin) is a small molecular weight antibiotic (Fig. 16) originally isolated from *Streptomyces venezuelae* in 1947.¹⁴⁷ The anti-phage effect of chloramphenicol was first reported in 1954 on *E. coli* phage T1, where bacteriostatic concentrations of chloramphenicol completely arrested phage multiplication in the host cells.¹⁴⁸ Subsequent studies showed that chloramphenicol is active against a wide panel of coliphages^{149–151} and *Streptococcus* phages.¹⁵² Chloramphenicol treatment was shown to not affect phage adsorption and DNA penetration¹⁵⁰ but to inhibit phage protein synthesis.¹⁵³ The protein synthesis inhibition was reversible (*i.e.*, it was relieved after removing chloramphenicol from phage-infected cells).¹⁵³ In some cases, chloramphenicol

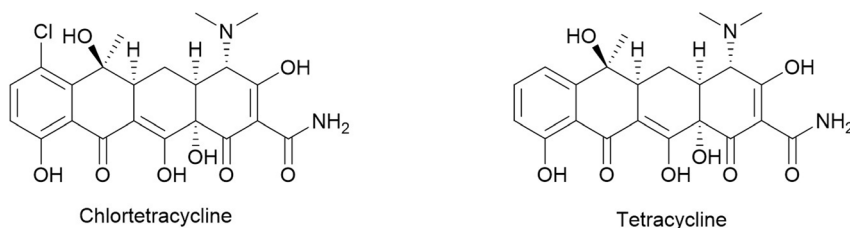


Fig. 15 Chemical structures of tetracyclines.



also inhibited phage DNA synthesis, which is presumably due to the indirect effect of peptide elongation inhibition,^{125,150,154–156}

In addition to the direct inhibitory action of chloramphenicol on phage protein synthesis, an earlier study reported that chloramphenicol-resistant bacteria also exhibited resistance to phages.¹⁵⁷ Chloramphenicol-resistant *Streptococcus* mutants, which evolved during chloramphenicol treatment, became resistant to phage infections. The mechanism of dual resistance to the antibiotic and phages was unclear. The authors found no evidence that the bacterial cell wall was modified in the mutants. In fact, the phages were able to adsorb and inject their genome into the mutants as well as they could into wild-type bacteria. The chloramphenicol-resistant ribosomes may somehow be immune to hijacking by phages.

Chloramphenicol treatments have also been shown to encourage the temperate coliphage P1 to enter its lysogenic phase, while the detailed mechanism is still unclear.¹⁵⁸

2.2.4. Erythromycin A. Erythromycin A is a macrolide antibiotic, comprising a 14-membered macrocyclic lactone with two sugar moieties attached (Fig. 17).¹⁵⁹ Erythromycin A was first isolated from *Saccharopolyspora erythraea* in 1952,¹⁶⁰ and later was shown to inhibit SPO1 phage multiplication in *B. subtilis* due to impeded phage protein synthesis.¹⁶¹ Erythromycin A interacts with host ribosomes and only allows the synthesis of short peptides with 6–8 amino acids before translation aborts (Fig. 14).^{125,154,155} As discussed above, the inhibitory action of erythromycin A on host ribosomes is likely the reason why phage protein synthesis is also inhibited during infection, as phage protein synthesis solely relies on host ribosomes. Slight inhibition of phage DNA synthesis was also observed upon erythromycin A treatment, possibly as a result of hampered synthesis of phage-encoded DNA replication machinery.¹⁶¹

2.2.5. Streptothricin. Streptothricins were among the first antibiotics discovered from soil Actinomycetota.¹⁶² Streptothricin F is the major component of an antibiotic mixture that was first isolated from *Streptomyces lavendulae* in 1942,¹⁶³ which features a streptolidine lactam ring, a glucosamine sugar, and a β -lysine (Fig. 18). Soon after its discovery, the anti-phage activity of streptothricin F was reported against two *E. coli* phages in 1945, where both plaque formation and phage

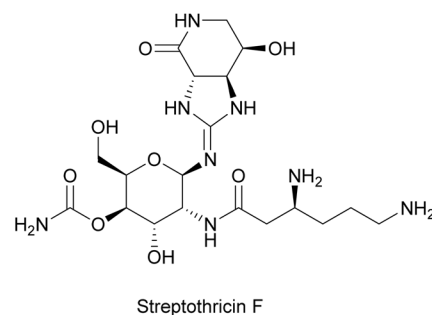


Fig. 18 Chemical structure of streptothricin F.

multiplication were inhibited.¹¹⁴ Subsequent work revealed inhibition of influenza virus, as well.¹⁶⁴ Streptothricin F is a protein synthesis inhibitor¹⁶⁵ that results in miscoding during peptide elongation¹⁶⁶ and impeded ribosomal translocation¹⁶⁷ (Fig. 14). The phage inhibition effect from streptothricin F is likely due to its inhibitory action on host ribosomes, thus interfering with the expression of essential phage proteins as discussed above.

2.2.6. Elfamycins. Elfamycins (Fig. 19) are a class of structurally diverse antibiotics that specifically target prokaryotic elongation factor thermo unstable (EF-Tu) during protein synthesis.^{168,169} In 1972, the first member of this class, kirromycin, was isolated from *Streptomyces collinus*.¹⁷⁰ Following the discovery of kirromycin, several of its analogs were also isolated, such as factumycin (A40A) from *Streptomyces lavendulae*,¹⁷¹ A73A from *Streptomyces viridifaciens*,¹⁷² and RK-1009 from *S. griseus*.¹⁷³ These analogs were shown to inhibit plaque formation by bacteriophage B on *S. griseus*,¹⁷³ presumably due to inhibition of phage protein synthesis steps that involve EF-Tu.¹⁷⁴ *N*-Methyl kirromycin (aurodox) in the elfamycin family has been shown to inhibit EF-Tu-assisted tRNA delivery (Fig. 14).¹⁷⁵ As factumycin, A73A, and RK-1009 share structural similarity with aurodox, they likely interfere with bacterial ribosomes in a similar manner, which eventually inhibits phage protein synthesis and hampers phage reproduction.

2.3. Dysregulate protein degradation (acyldepsipeptides)

Acyldepsipeptides (Fig. 20) are a relatively new class of antibiotics with promising results against multidrug-resistant pathogens because of their distinct mechanism of action.^{176–178} The first acyldepsipeptide, A54556A, was isolated from *Streptomyces hawaiiensis* in 1985.¹⁷⁹ Six years later, another member in this class, enopeptin A, was isolated in a screen for anti-phage natural products.¹⁸⁰ It was shown that enopeptin A produced by *Streptomyces* sp. RK-1051 inhibited plaque formation from bacteriophage B on *S. griseus*.¹⁸⁰

It is still unclear how enopeptin A inhibits phage proliferation, but it is likely due to the dysregulation of host proteolytic systems. Regulated proteolysis maintains a healthy proteome by identifying and degrading damaged and unneeded proteins.¹⁸¹ The caseinolytic protease (Clp) complex is one of the main proteolytic systems in bacteria.¹⁸² In the Clp complex, 14 units of protease ClpP form a proteolytic chamber, whose activity is

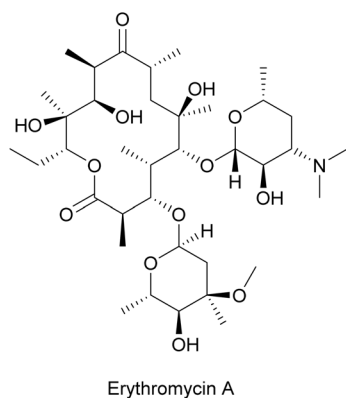


Fig. 17 Chemical structure of erythromycin A.



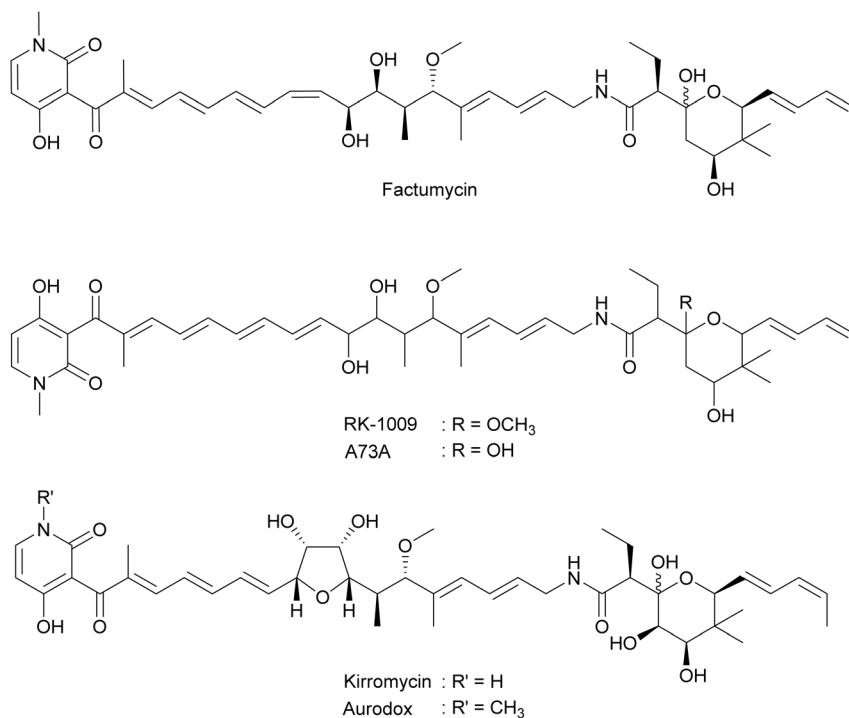


Fig. 19 Chemical structures of elfamycins.

regulated by ATPase ClpX/A/C, which recognizes damaged proteins, unfolds them, and threads them into the proteolytic chamber.¹⁸³ Acyldepsipeptides bind at the ATPase pocket on the

ClpP surface and dysregulate the proteolytic activity of ClpP complex.¹⁸⁴ Dysregulation of the host proteolytic system by acyldepsipeptides could be detrimental to the phage infection

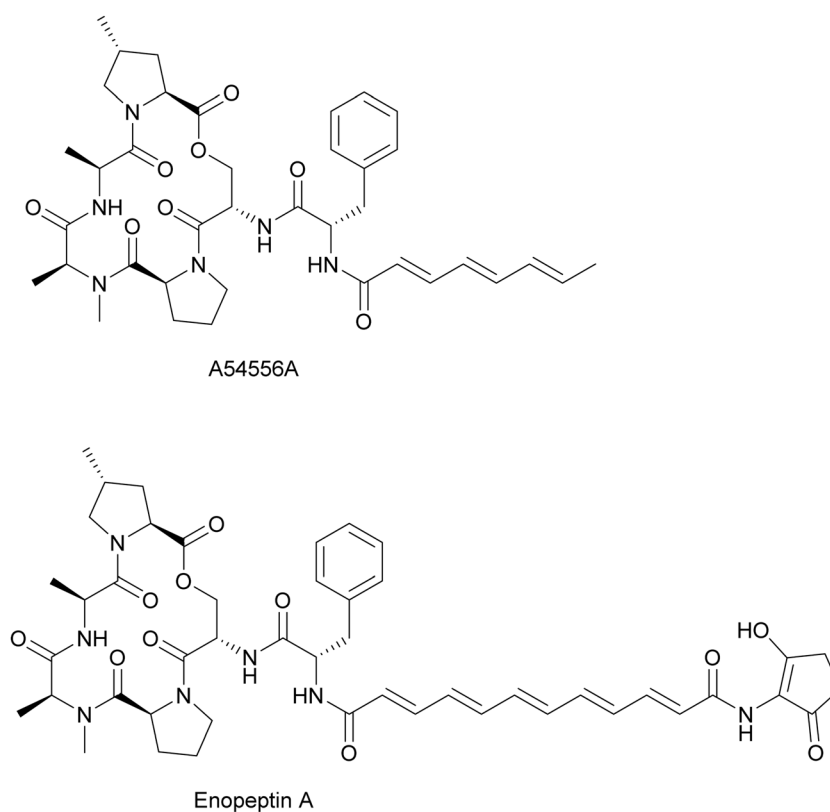


Fig. 20 Chemical structures of acyldepsipeptides.



process in two ways. First, the uncontrolled degradation of nascent peptides could prevent the production of phage proteins. Additionally, some phages encode proteins that regulate host proteolytic systems like the Clp complex. By regulating proteolysis, phages can optimize amino acid flux for their own development.¹⁸⁵ Therefore, inhibition of Clp may also inhibit phage proliferation by preventing this phage-based reprogramming of proteolysis.

2.4. Sequester iron (siderophores)

Siderophores are structurally diverse microbial secondary metabolites with high affinity to iron.¹⁸⁶ These metabolites are synthesized and secreted by microbes to harvest iron from the environment, and then the iron-bound siderophores are transported back into the cells by specific transporters on the membrane.¹⁸⁶ Microbial species compete with each other for scarce environmental iron by making structurally distinct siderophores.¹⁸⁷ Due to the specificity of siderophore transporters, a siderophore made by one species often cannot be utilized by another species, thus sequestering iron away from competitors.¹⁸⁷

Recently, it has been reported that the *E. coli* siderophores enterobactin and linear enterobactin (Fig. 21A) can repress ICP1 phage proliferation in *Vibrio cholerae* by iron sequestration.¹⁸⁸ Enterobactin has a very narrow effective range, as it causes a complete growth arrest of *V. cholerae* at concentrations higher than 4 μM . In contrast, linear enterobactin is effective against phages over a wider range of concentrations, because it does not strongly inhibit *V. cholerae* growth even at 200 μM . This special trait of linear enterobactin is likely because *V. cholerae* can pirate linear enterobactin but not enterobactin for its iron

uptake.¹⁸⁹ Therefore, linear enterobactin probably induces a slight iron starvation in *V. cholerae* without completely arresting its growth. This modest iron deficiency in the host appears to inhibit active phage reproduction by delaying phage-mediated cell lysis and reducing the number of new phages produced by each infected cell.¹⁸⁸

The exact mechanism by which iron deficiency hampers phage proliferation is unclear. Iron is necessary for many cellular processes, and it is plausible that an iron deficiency inhibits several processes that are more essential for phage replication than for host replication.¹⁸⁸ One hypothesis is that iron deficiency in the host compromises the activity of phage-encoded ribonucleotide reductases (RNRs), thus repressing phage DNA synthesis. RNR is a vital enzyme for DNA synthesis. It converts nucleotides into deoxynucleotides (Fig. 21B).¹⁹⁰ Phage-encoded RNRs are important for rapid phage DNA synthesis¹⁹¹ and effective reproduction.¹⁹² ICP1 phage encodes a class Ia RNR and a class III RNR on its genome,¹⁹³ both of which require iron as a cofactor.¹⁹⁰ It has been shown that intracellular iron deficiency caused by an iron chelator can attenuate RNR activity in human cells.¹⁹⁴ Therefore, it is possible that enterobactin and linear enterobactin sequester iron away from ICP1-infected *V. cholerae*, which inhibits ICP1-encoded RNR activity and impedes rapid ICP1 proliferation. Further experiments are required to distinguish this mechanism from the numerous other influences of iron starvation.

2.5. Modify or down-regulate phage receptors

In contrast to the previously discussed anti-phage natural products that interfere with phage reproduction within the host

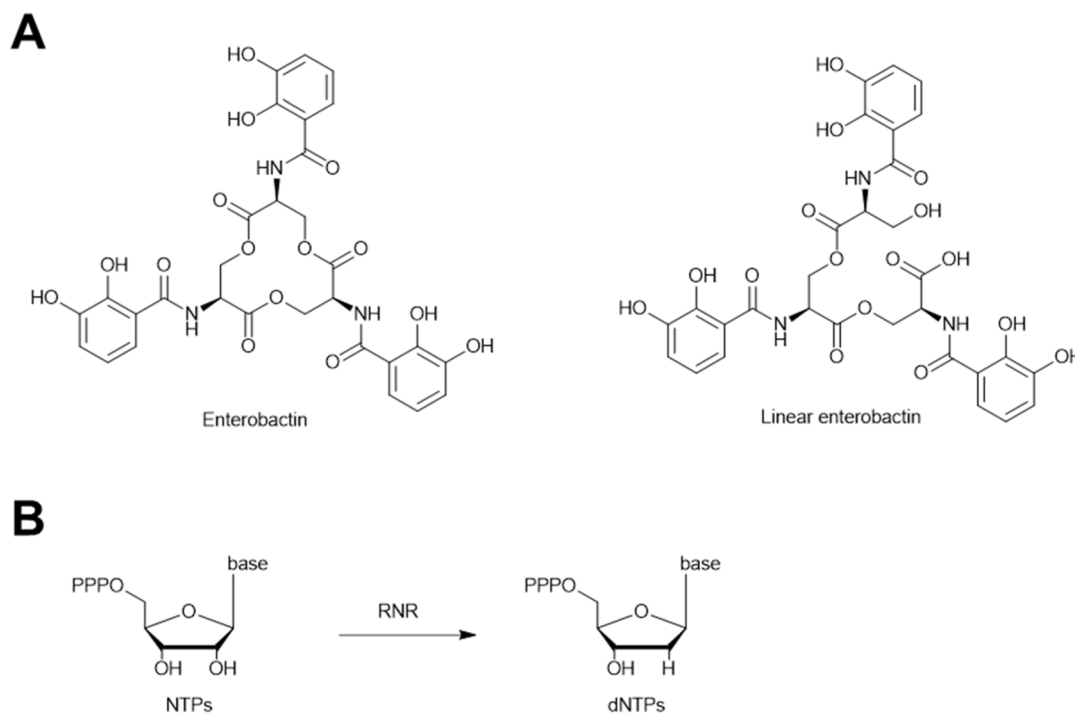


Fig. 21 Siderophores inhibit phage infections. (A) Chemical structures of anti-phage siderophores. (B) The reaction catalyzed by ribonucleotide reductase (RNR).



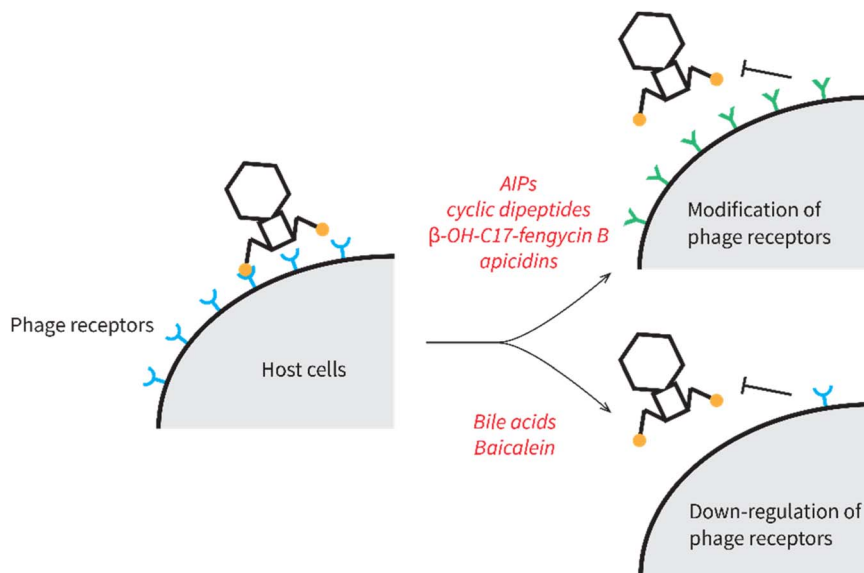


Fig. 22 Mechanisms of action of natural products that influence phage adsorption.

cell, others inhibit the initial adsorption of phages to their host surfaces (Fig. 22). Reduced adsorption is mediated by modifications to the bacteria cell surface receptors that phages recognize for binding and infection. These receptors can be modified either qualitatively by changing their composition or quantitatively by decreasing their expression level.

2.5.1. Autoinducing cyclic peptides (AIPs). Autoinducing cyclic peptides (AIPs) are chemical signals produced by *Staphylococcus* bacteria that encode the *agr* quorum sensing (QS)

system (Fig. 23A).¹⁹⁵ The *agr* QS system coordinates group behaviors of *Staphylococcus* in response to various AIP concentrations at different cell densities.^{196,197} There are four variants (I–IV) of the *agr* system in *S. aureus*, and each variant of the *agr* system is only induced by its cognate AIP. In fact, non-cognate AIPs produced by other bacteria often inhibit the natural functioning of an *agr* system.¹⁹⁷ Recently, it was shown that cognate AIP-I can promote Stab20 phage infections in *S. aureus* encoding *agr*-I, while the non-cognate AIP-hy produced by *Staphylococcus*

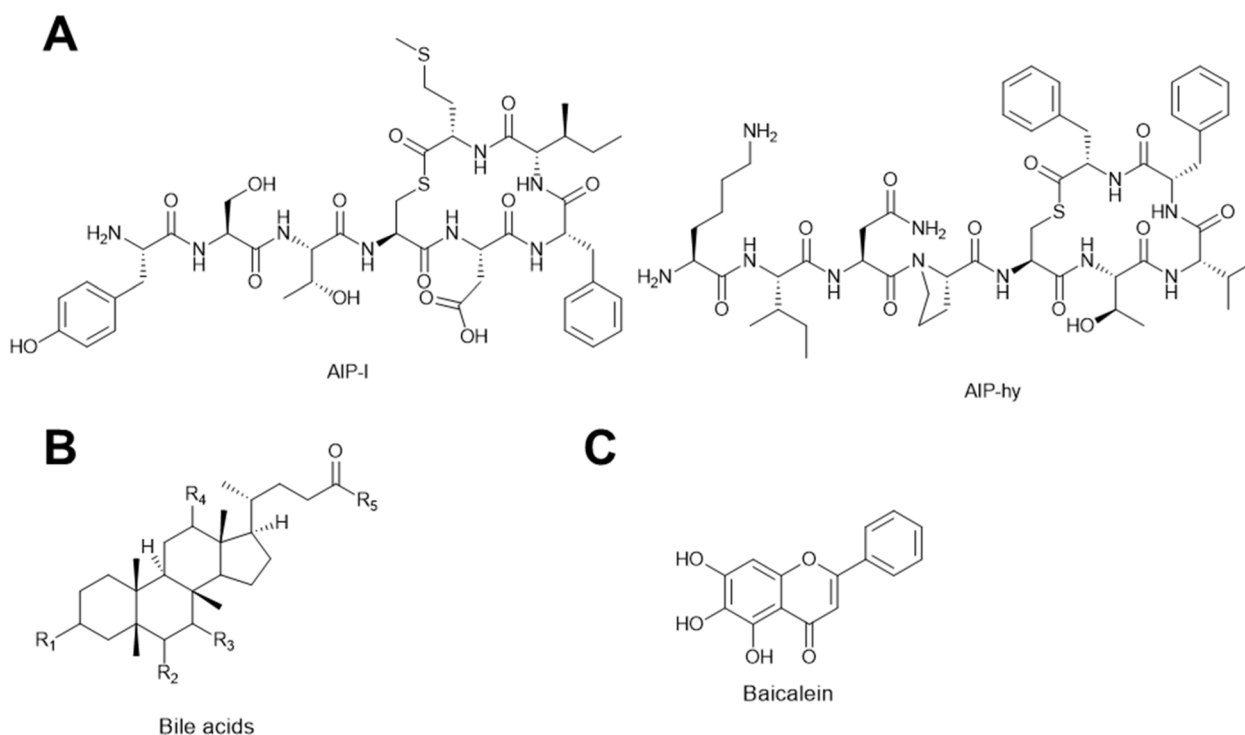


Fig. 23 Natural products that modulate or down-regulate phage receptors on the host cell surface. (A) Chemical structures of *agr* inhibitors. (B) Generic chemical structure of bile acids. (C) Chemical structure of baicalein.



hyicus inhibits phage infections in *S. aureus* encoding *agr-I*, as measured by plaque formation and phage-induced host cell lysis.¹⁹⁸ The AIPs were shown to influence phage infectivity by modifying the phage receptor on the cell surface. Namely, the AIPs changed the expression levels of *tarM*. TarM is an enzyme that adds α -N-acetylglucosamine to the wall teichoic acid (WTA),¹⁹⁹ which blocks Stab20 phage adsorption.¹⁹⁸ The cognate AIP-I induces *agr-I*, which represses *tarM* expression, thus facilitating phage adsorption. On the contrary, noncognate AIP-hy inhibits *agr-I* activation, thereby derepressing *tarM* and inhibiting phage adsorption. Furthermore, by co-culture assays, the authors discovered that other *Staphylococcus* strains that frequently co-occur with *S. aureus* on the skin of humans and animals also exhibit an anti-phage effect on *S. aureus*, presumably through secretion of inhibitory non-cognate AIPs.¹⁹⁸ Therefore, cross-species metabolic interactions can dramatically impact phage infection outcomes in *Staphylococcus*.

2.5.2. Bile acids. Bile acids (Fig. 23B) are a group of cholesterol metabolites with important functions in vertebrate animals, such as facilitating lipid absorption and excretion in the gut, lipid and glucose homeostasis, and immune signaling.²⁰⁰ Bile acids are synthesized from cholesterol in the liver as primary bile acids, which are further metabolized by the gut microbiota into secondary bile acids. It was recently discovered that *V. cholerae*, a gut pathogen, became resistant to ICP1 phage infections when exposed to host bile acids.²⁰¹ The authors showed that the phage resistance occurred due to a transient modification of phage receptors on the cell wall in response to a combination of bile acids, anaerobicity, and low pH.²⁰¹ The O-antigen (or outer core polysaccharide) of lipopolysaccharide on the *V. cholerae* cell wall is the receptor of ICP1 phages.²⁰² When exposed to bile acids at a low pH under anaerobic conditions, O-antigen synthesis in *V. cholerae* was impaired due to a decrease in O-antigen biosynthetic enzyme levels and a depletion of central carbon metabolites required for constructing O-antigen.²⁰¹ As a result, the decreased O-antigen on the cell surfaces inhibited ICP1 adsorption, thus limiting its infection. This effect may synergize with the aforementioned influence of iron sequestration¹⁸⁸ to explain transient phage-resistance observed in a prophylaxis phage therapy experiment against *V. cholerae* conducted in animal guts.²⁰³

2.5.3. Baicalein. Baicalein is a flavonoid compound originally isolated from the roots of *Scutellaria baicalensis* (Fig. 23C).²⁰⁴ Recently, it has been shown to inhibit DMS3vir phage infection in *Pseudomonas aeruginosa* through phage adsorption inhibition.²⁰⁵ DMS3vir phage requires the type IV pilus of *P. aeruginosa* as its receptor,^{206,207} which is regulated by quorum sensing systems.²⁰⁸ The authors proposed that the inhibited phage adsorption was due to the down-regulation of type IV pilus through the inhibition of quorum sensing caused by baicalein.²⁰⁵

2.6. Activating anti-phage defense systems

Over millennia of co-evolution between bacteria and phages, bacteria have acquired hundreds of anti-phage defense systems to provide protection against phage infection.²⁰⁹ Most of these systems were cryptic genes within bacterial genomes for

decades until recent advances in bioinformatic analysis revealed the mystery of these prokaryotic “immune systems”. A large fraction of these systems rely on nucleotide-derived signaling molecules to abort phage infections,²¹⁰ such as CBASS,^{211,212} Thoeris,^{213–215} type III CRISPR,^{216–218} and Pycsar.²¹⁹ Cumulatively, systems of this type are present in ~36% of sequenced bacterial genomes.²¹⁰ Generally, these defense systems utilize a sensor protein to sense phage infection and convert cellular nucleotides into secondary signaling molecules. These “immune signals” then bind and activate downstream effector proteins to abort phage infections. CBASS has the most diverse signal molecules among the immune signaling systems, with more than 10 distinct nucleotide signals identified so far. CBASS signal molecules feature cyclic di- or trinucleotide species, with combinations of both purine and pyrimidine bases that are linked through 3′–5′ and/or 2′–5′ phosphodiester bonds (Fig. 24A). Thoeris systems have three types of signals identified so far, which are all derived from cellular NAD⁺ (Fig. 24B). In type III CRISPR systems, two types of signals have been discovered, including cyclic oligoadenylate and SAM-AMP (Fig. 24C). Pycsar systems exclusively synthesize cyclic pyrimidine mononucleotides as signal molecules, such as 3′,5′-cyclic cytosine monophosphate (cCMP) and 3′,5′-cyclic uridine monophosphate (cUMP) (Fig. 24D).

The immune signaling molecules are unique and distinct from other anti-phage natural products in multiple ways. First, although immune signals are specialized to antagonize phage activity, they mostly activate toxic effectors that lead to cell death before phage infection completes.²¹⁰ In contrast, many anti-phage natural products exhibit weak or no toxicity to the host cell at concentrations that inhibit phage infections. Second, the anti-phage action of immune signals has only been demonstrated in a cell-autonomous way. In other words, the signal from an infected cell does not activate defenses in neighboring cells. It would be interesting to investigate if immune signals can also act in non-cell-autonomous way and activate the anti-phage systems in the whole bacterial community to confer “herd immunity” and to shape microbial ecology. Third, the diverse anti-phage immune signaling systems provide an opportunity for systematic discovery of anti-phage molecules, since the signals produced by many of the systems remain unknown.

3. Phage-promoting natural products

Natural products that promote phage infections have been reported less than those that inhibit phages. However, a revived interest in phage therapy has motivated the discovery of natural products (especially antibiotics^{16–19}) that can synergize with phages for antibacterial therapy. Beyond their therapeutic potential, phage-promoting natural products might also play an important role in mediating microbial competition in nature. For example, phage-promoting metabolites produced by one microbe could sensitize its competitors to phage predation, thus giving the producer a competitive advantage over other bacteria.²²⁰

The phage-promoting activity of natural products can be assessed experimentally in multiple ways. We highlight two



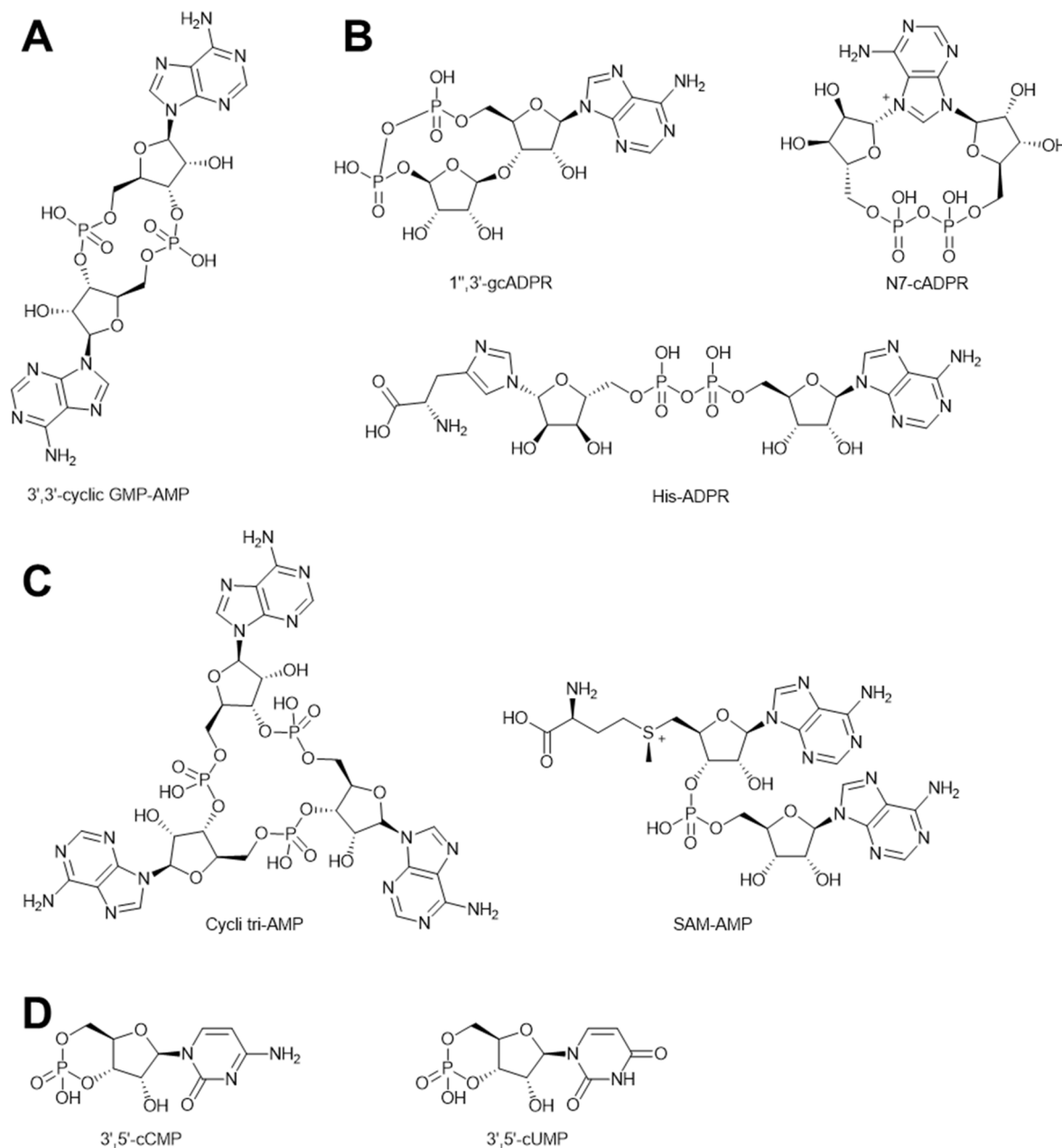


Fig. 24 Natural products that activate anti-phage immune systems (A) chemical structure of a representative CBASS signal. (B) Chemical structures of Thoiris signals. (C) Chemical structure of representative type III CRISPR signals. (D) Chemical structures of Pycsar signals.

methods (Fig. 25) that can reveal selective phage lysis promotion by molecules that are not antimicrobial (at least at the applied dose). One case monitors the increase in area of plaques (areas of phage-induced bacterial lysis on an agar surface). The other case monitors improved phage-induced lysis in liquid culture.

The current known natural products that promote phage infections are discussed in the following sections according to their specific mechanisms.

3.1. Inhibit peptidoglycan synthesis (beta-lactams)

β -Lactam compounds are arguably the most widely prescribed antibiotics, representing more than half of all commercially available antibiotics in use.²²¹ This family of antibiotics is

named after their shared feature: a β -lactam ring (Fig. 26A). In 1929, penicillin was isolated by Alexander Fleming²²² from *Penicillium rubens*.²²³ In 1947, penicillin was first reported to accelerate phage-mediated host cell lysis in *Staphylococcus aureus*.²²⁴ Many other classes of β -lactam antibiotics (Fig. 26A) have been isolated from microbes and further expanded by medicinal chemistry efforts,²²⁵ such as cephalosporins,²²⁶ carbapenems,²²⁷ and monobactams.²²⁸ All of these β -lactam antibiotics synergize with phages to kill a variety of bacterial hosts.^{16,229}

The synergy between β -lactam antibiotics and phages is presumably due to the impeded cell wall synthesis caused by β -lactams.^{16,230} One of the key structures of bacterial cell walls is



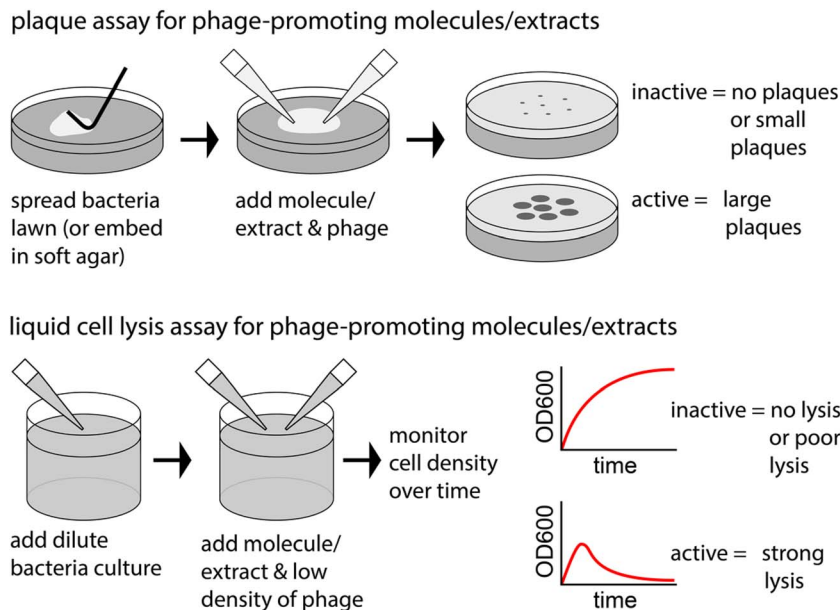


Fig. 25 Example experiments to observe phage-promoting natural products.

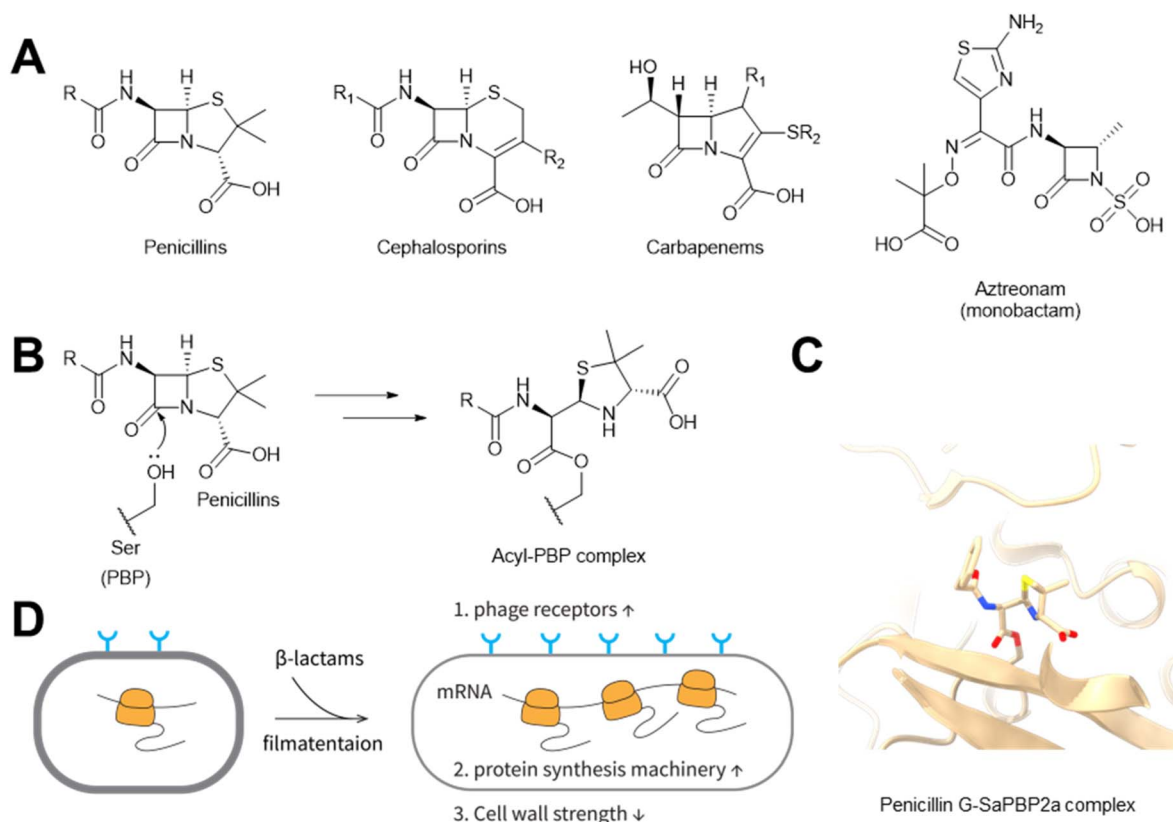


Fig. 26 β -Lactams promote phage infection by inhibiting peptidoglycan synthesis. (A) Chemical structures of β -lactams. (B) Formation of an acyl-enzyme complex between β -lactams and PBPs. (C) Structure of penicillin G in complex with PBP2a from *S. aureus*. [PDB: 1MWT]. (D) Possible mechanisms by which filamented cells promote rapid phage proliferation.

peptidoglycan, whose synthesis is catalyzed by penicillin-binding proteins (PBPs).^{231,232} β -Lactams can occupy the active site of PBPs and form covalent acyl-enzyme complexes that

arrest PBP-catalyzed peptidoglycan cross-linking (Fig. 26B and C).^{233–235} At sublethal concentrations of β -lactams, the hampered peptidoglycan synthesis leads to poor cell division and filament



formation (Fig. 26D).^{16,236} On one hand, the filamented bacterial cells possess a larger cell surface, which facilitates the phage adsorption step.²³⁰ On the other hand, the inhibited cell division might cause each bacterial filament “cell” to have more protein synthesis machinery, leading to a larger burst size (the number of new phages produced by each infected cell).^{16,230} Lastly, interrupted peptidoglycan synthesis may also ease the effort of peptidoglycan degradation by endolysins in the phage-mediated cell lysis step, which would expedite cell lysis.¹⁶

3.2. Inhibit stationary phase transition

Transitioning into stationary phase and ultimately cell dormancy are common strategies for bacteria to adapt to environmental stresses.²³⁷ This transition can further afford

recalcitrance to phage infection. For example, in *Bacillus*, multiple pathways regulated by Spo0A during stationary phase transition can repress phage activities (Fig. 27A).²²⁰ In the dormant state, the altered cell wall²³⁸ and heavily reduced metabolic activity²³⁷ can block phage adsorption²³⁸ and inhibit rapid phage proliferation,^{239,240} respectively (Fig. 27A). Therefore, molecules that inhibit the stationary phase transition and cell dormancy could keep bacterial hosts in their phage-sensitive states, thus promoting phage reproduction. Three examples of natural products with this ability follow.

3.2.1. Siderophores. A recent study showed that a siderophore produced by *Streptomyces*, coelichelin (Fig. 27B), can inhibit the stationary phase transition in *B. subtilis*, thus promoting phage predation on *B. subtilis*.²²⁰ Iron sequestration caused by coelichelin can block the activation of Spo0A,²²⁰ the

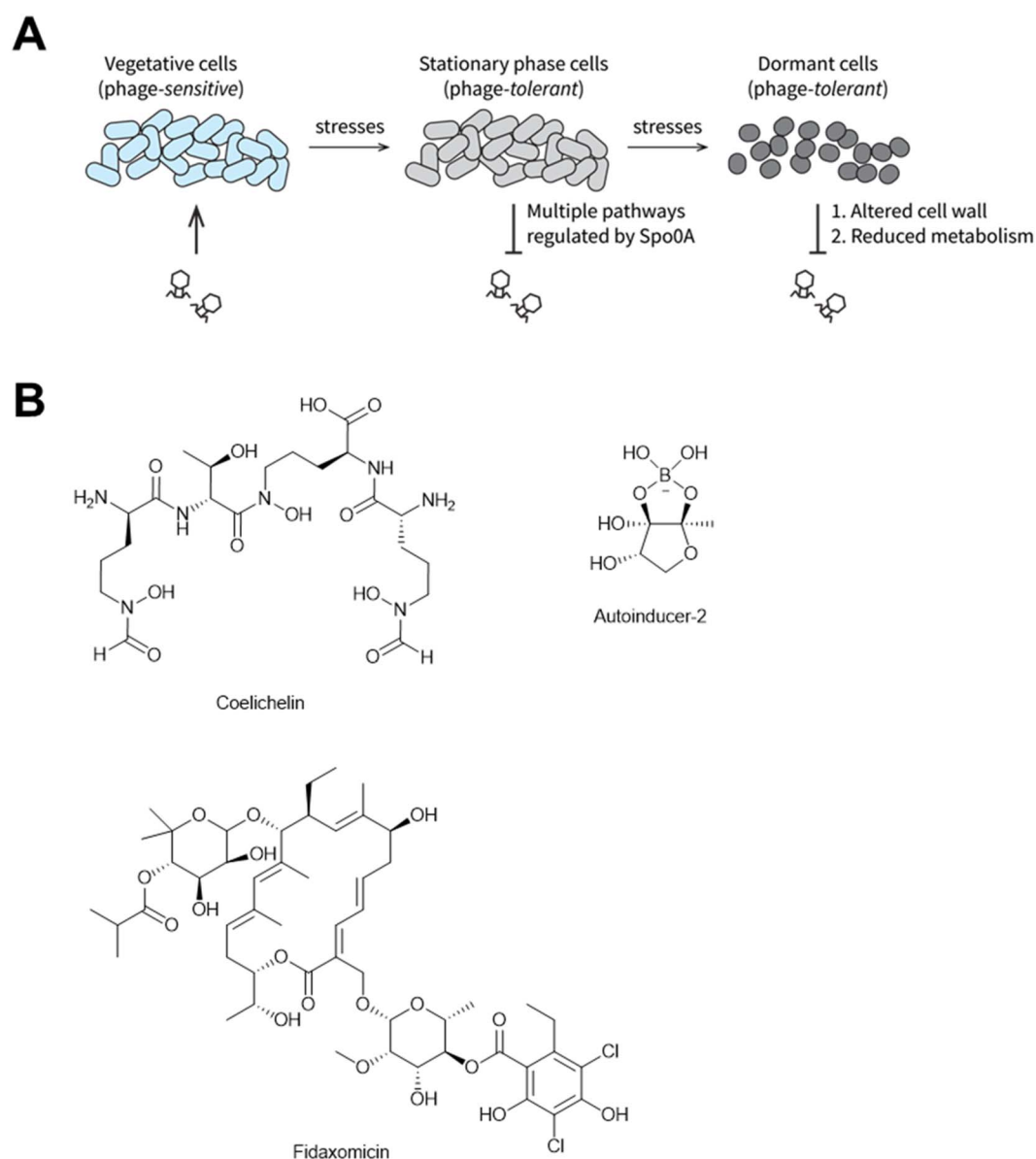


Fig. 27 Natural products that promote phage infections by inhibiting stationary phase transition. (A) Mechanisms by which stationary phase transition and dormancy inhibit phage infection. (B) Chemical structures of Spo0A inhibitors.



master transcriptional regulator in *B. subtilis* that controls the transition to stationary phase.^{241,242} The authors further showed that coelichelin production gave *Streptomyces* a competitive advantage over *B. subtilis* by sensitizing *B. subtilis* to phage infection.²²⁰ They found that other siderophores (and even non-

siderophore metabolites) also exhibited similar phage-promoting activities.²²⁰

3.2.2. Other Spo0A inhibitors. Non-siderophore bacterial secondary metabolites have also been shown to inhibit Spo0A activation or expression. Namely, autoinducer-2 (ref. 243) and fidaxomicin (Fig. 27B)²⁴⁴ inhibit Spo0A activation in *Bacillus*

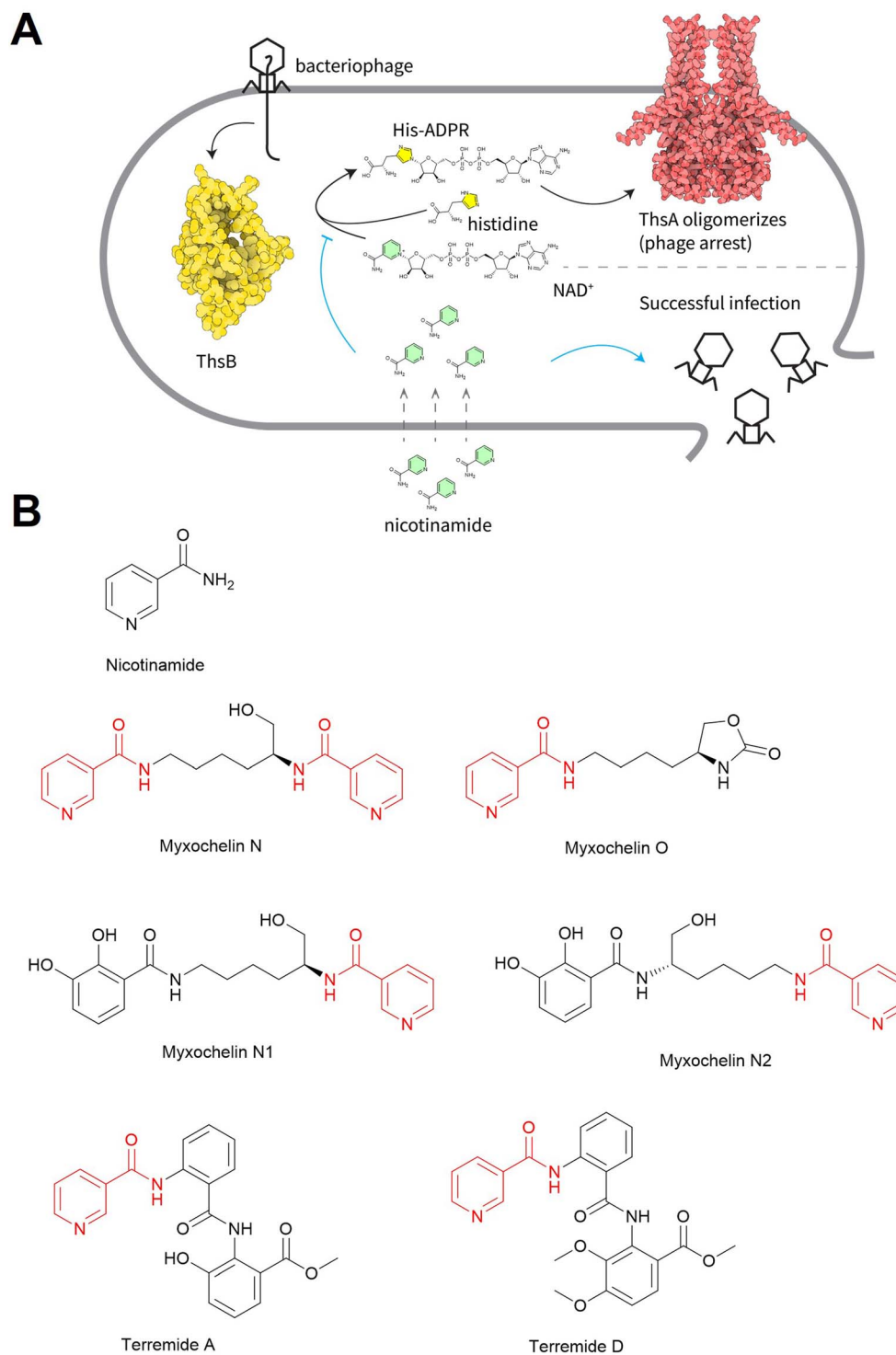


Fig. 28 Natural products promote phage infections by inhibiting the type II Thois anti-phage system. (A) Mechanism of action of the type II Thois system, which can be inhibited by exogenous nicotinamide. (B) Chemical structures of nicotinamide and nicotinamide-containing natural products.



velezensis and *Clostridioides difficile*, respectively. These Spo0A inhibitors have not been evaluated for their interaction with phages, but hypothetically they could promote phage infection.

Spo0A-regulated dormancy behaviors are found in many bacteria in the Bacillota (Firmicutes) phylum.²⁴⁵ The discoveries above suggest that inhibiting the Spo0A-mediated stationary phase transition and sensitizing competitors to phage predation could be a common competition strategy among microbes. It is likely that other natural Spo0A inhibitors exist and remain to be discovered.

3.3. Inhibit anti-phage defense systems

As discussed in the previous section, anti-phage immune signaling systems encode protein components that generate or bind small molecule signals.²¹⁰ These components possess cavities for small molecule binding, which could be targets for inhibition or activation by exogenous natural products.

Recently, it was reported that nicotinamide can inhibit the type II Thoeris antiphage system encoded in a wide range of

hosts, including *Bacillus amyloliquefaciens*, *P. aeruginosa*, and *Enterococcus faecalis*.²⁴⁶ In doing so, it promoted phage predation on these hosts. The type II Thoeris system relies on two proteins, ThsA and ThsB (Fig. 28A).²¹⁴ The ThsB protein can sense phage infection and generate a small molecule alarm signal, histidine-ADP-ribose (His-ADPR). The His-ADPR signal then activates ThsA, which arrests phage replication. Since the first step of His-ADPR biosynthesis is NAD⁺ hydrolysis into nicotinamide and ADPR by the TIR domain of ThsB, excess nicotinamide (Fig. 28A) inhibits NAD⁺ hydrolysis. Therefore, high concentrations of exogenous nicotinamide blocked His-ADPR production and restored phage infectivity.²⁴⁶ Beyond nicotinamide, some microbes also produce nicotinamide-containing secondary metabolites, such as myxochelins (Fig. 28B)^{247,248} and terremides (Fig. 28B),^{249,250} which may also inhibit the type II Thoeris system through a similar mechanism of action. Although yet to be demonstrated, nicotinamide and its analogs may also inhibit other immune systems that contain TIR domains.

This discovery is the first demonstration that anti-phage systems can be inhibited by small molecule natural products. Considering the presence of dozens of anti-phage systems, we anticipate that natural products targeting other systems exist and remain to be discovered.

3.4. Unknown mechanism (rebaudioside A)

Rebaudioside A is a natural high-potency sweetener isolated from stevia leaves (Fig. 29).²⁵¹ Recently, it was found that rebaudioside A facilitated phage infection on *Yersinia enterocolitica*.²⁵² Preliminary data suggested that phage adsorption was promoted by rebaudioside A treatment.²⁵² The increased adsorption might be due to a stabilizing effect imposed on the free phage particles by rebaudioside A.²⁵² The phage particles might aggregate in suspensions. Rebaudioside A may prevent phage aggregation,

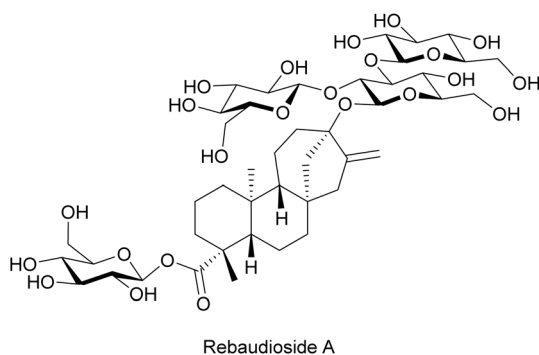


Fig. 29 Chemical structures of rebaudioside A.

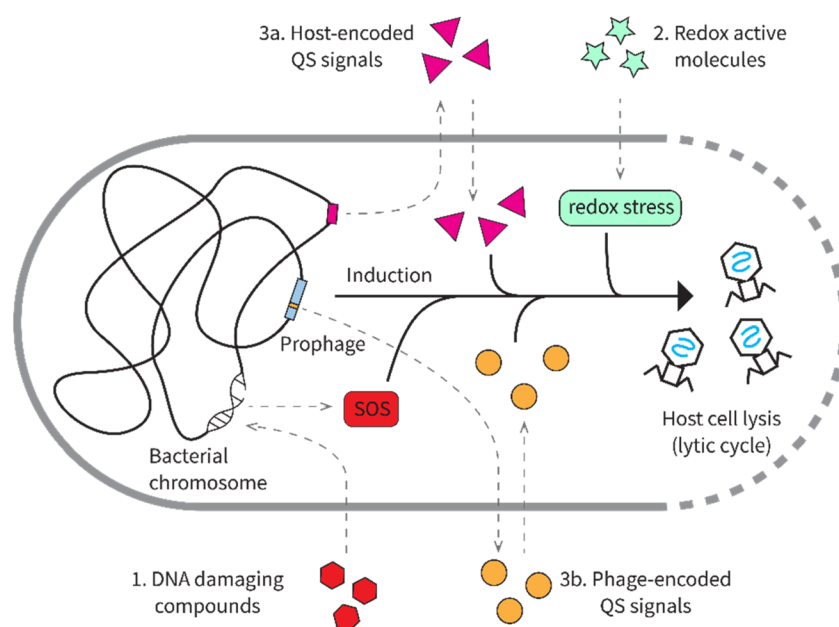


Fig. 30 The four major mechanisms of prophage induction.



which increases the effective phage titer.²⁵² The validation of this hypothesis and the exact mechanism of action of rebaudioside A still require further investigation in the future.

4. Lysis/lysogeny-regulating natural products

In contrast to lytic phages that exclusively undergo lytic cycles, temperate phages can undergo both the lytic cycle and the lysogenic cycle.^{253,254} In the lysogenic life cycle, temperate phages integrate their genomes into the chromosome of their host bacteria. Here they lay dormant as “prophages”, replicating along with the host genome and propagating into all of the

progeny of that host cell.^{253,254} Temperate phages can then switch back to their lytic life cycle when conditions would benefit lysis—either in response to environmental signals like microbial metabolites or through phage-encoded quorum sensing systems (Fig. 30). In this section, we will review the known microbial metabolites that regulate lysis-lysogeny “decisions” in temperate phages.

4.1. Damage DNA

One mechanism that induces many prophages to enter their lytic cycle is DNA damage in the host (Fig. 30). This behavior provides a clear fitness benefit to the prophage. Host cells undergoing excessive DNA damage will likely fail to replicate,

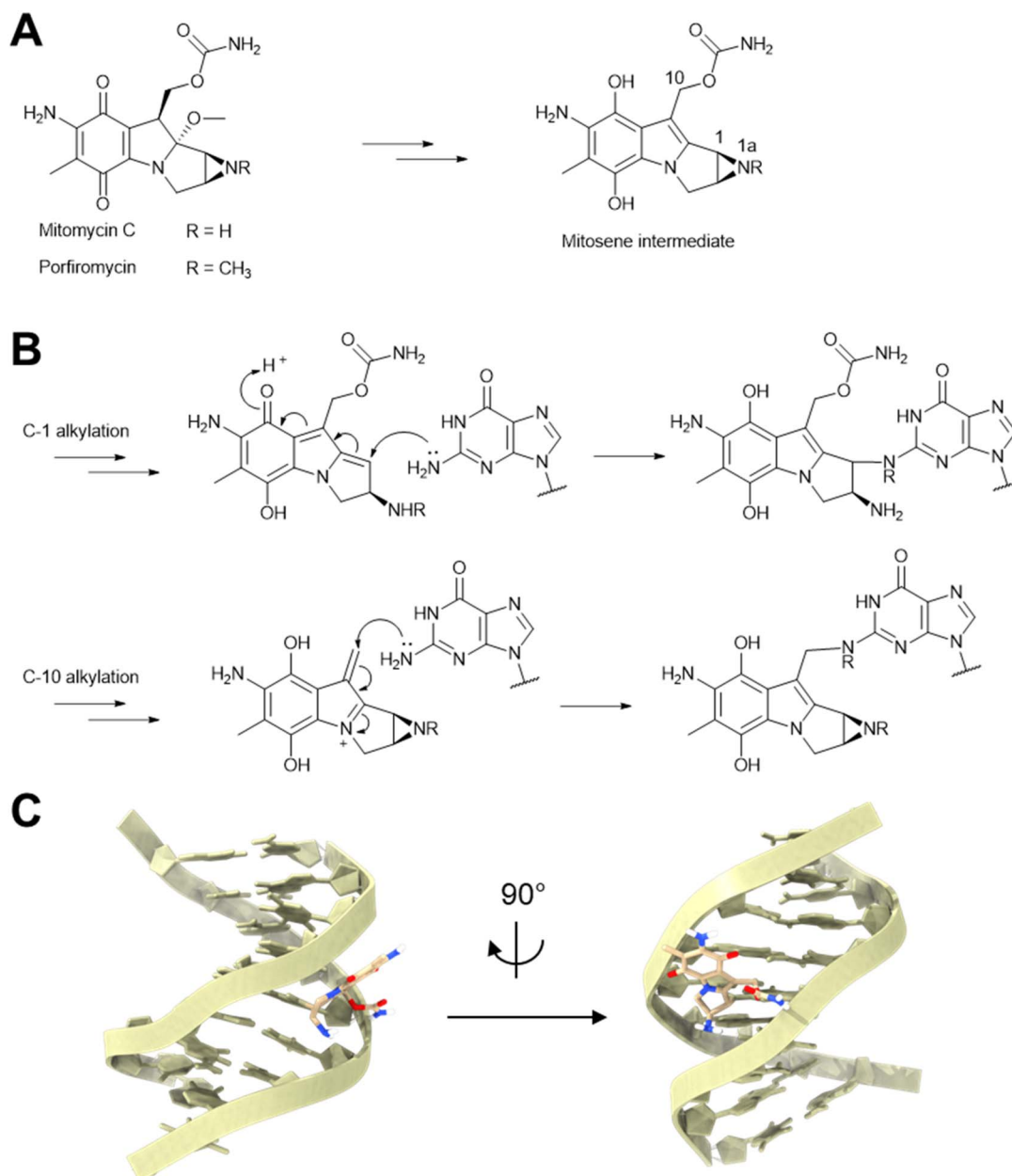


Fig. 31 The prophage induction mechanism of mitomycins. (A) Chemical structures of mitomycins and reactive mitosene intermediates. (B) Mechanism of DNA alkylation by mitomycins. (C) Structure of mitomycin C-DNA adduct through C1 alkylation [PDB: 199D].



which would arrest prophage propagation. Therefore, the phage should benefit from switching to the lytic cycle, where it could disperse dozens of phage particles to infect healthy cells. Since DNA-damaging agents have been attractive lead molecules for antitumor drugs, many prophage-inducing natural products were actually discovered in screens for antitumor drugs using *E. coli* containing the λ prophage.^{13–15} For λ and many other temperate phages, the lysogenic state is maintained by repressor proteins that prevent the transcription of lytic genes.²⁵⁵ DNA damaging natural products trigger the SOS response in the host bacterial cells.^{256–258} The activated SOS pathway in the host typically derepresses the lytic genes through one of two mechanisms: (1) autoproteolysis of the repressor proteins in a RecA-dependent manner²⁵⁹ or (2) expression of antirepressor proteins that antagonize repressor proteins.²⁶⁰

4.1.1. Mitomycins. Mitomycins are a family of potent antibiotics and antitumor drugs, composed of aziridine, quinone, and carbamate moieties arranged on the pyrrolo[1,2-*a*] indole core structure (Fig. 31A).²⁶¹ In 1958, mitomycin C was first isolated from *Streptomyces caespitosus*.²⁶² One year later, it was found that mitomycin C could induce the λ prophage in *E. coli* to enter its lytic cycle.²⁶³ Subsequently, mitomycin C treatment has become a standard protocol for prophage induction. Following the discovery of mitomycin C, the *N*-1a-methyl derivative porfiromycin isolated from *Streptomyces ardens*,²⁶⁴ was also shown to induce the lytic cycle of the λ prophage.¹⁴

The prophage induction capability of mitomycins can be attributed to their ability to alkylate DNA. Following activation through an enzymatic or chemical reduction pathway, mitomycins are converted into reactive mitosene intermediates (Fig. 31A).^{265,266} In the mitosene intermediate, electrophilic centers can be formed at either the C-1 or C-10 position and react with N-2 of guanine (Fig. 31B and C), generating either inter- or intra-strand DNA crosslinks.²⁶⁷ The DNA crosslinks activate the SOS response in the host cell,²⁶⁸ thus leading to prophage induction.

4.1.2. Azaserine. Azaserine is a naturally occurring derivative of serine with an α -diazooester moiety. It exhibits antibiotic and anti-cancer properties (Fig. 32A).²⁶⁹ Azaserine was first isolated from *Streptomyces fragilis* in 1954.^{270,271} Shortly following its discovery, azaserine was found to induce λ prophage in *E. coli*.^{14,272} As with mitomycin, the prophage induction activity of azaserine is presumably through a DNA alkylating mechanism. The diazo group in azaserine can undergo protonation to generate the diazonium moiety, which readily decomposes into a carbonium that can alkylate DNA (Fig. 32A).^{273,274} It has been reported that azaserine mainly reacts with purines, and subsequent spontaneous hydrolysis and/or decarboxylation forms *N*⁷-carboxymethylguanine, *O*⁶-carboxymethylguanine, or *O*⁶-methylguanine (Fig. 32B).^{275,276} DNA alkylation by azaserine has been reported to cause extensive DNA damage in bacterial hosts,^{277,278} which subsequently triggers the SOS response.^{277,278} This SOS response likely induces the lytic cycle in a similar manner as above.

4.1.3. Pluramycin A. Pluramycin A was first isolated from *Streptomyces pluricolorscens* in 1956 (Fig. 33).²⁷⁹ The prophage induction activity of pluramycin A was reported in many studies in the 1960s using λ prophage-containing *E. coli* strains.^{13,14,280} Pluramycin A is structurally similar to the earlier discussed molecule neopluramycin (Fig. 4). Like neopluramycin, pluramycin A is also capable of intercalating DNA.⁵⁰ Notably, the presence of an epoxide ring in pluramycin A (Fig. 33) allows it to react with the N-7 in the guanine base (Fig. 33).²⁸¹ DNA alkylation caused by pluramycin A may induce cellular DNA damage in bacterial hosts²⁸² and a subsequent SOS response, therefore inducing prophages to switch to the lytic cycle as discussed above.

4.1.4. Streptozotocin. Streptozotocin was first isolated from *Streptomyces achromogenes* in 1957 as an antibiotic (Fig. 34A).²⁸³ Later streptozotocin was shown to induce λ prophage in *E. coli*.¹⁴ The nitrosourea group in streptozotocin spontaneously decomposes into a diazene hydroxide (Fig. 34B).²⁸⁴ Specifically,

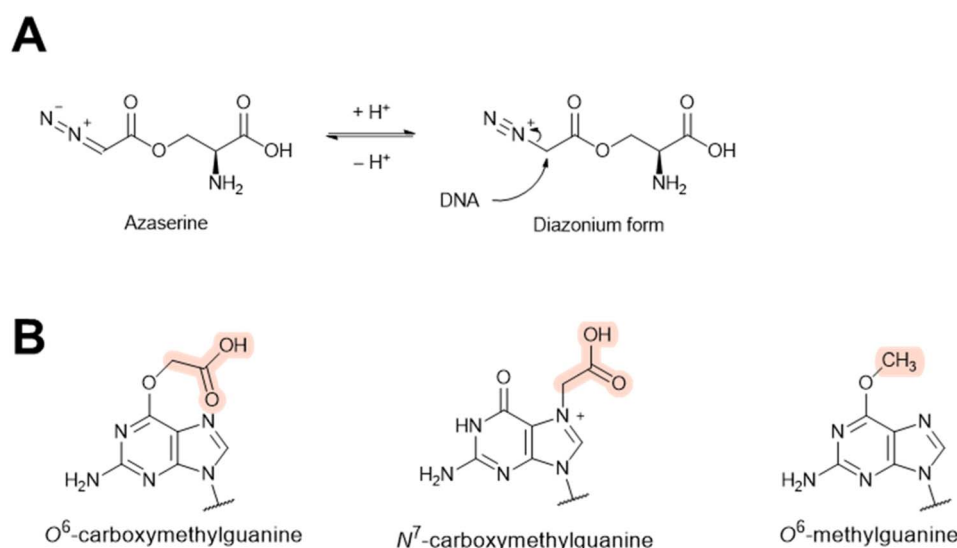


Fig. 32 The prophage induction mechanism of azaserine. (A) DNA alkylation mechanism of azaserine. (B) Three possible purine modifications by azaserine (highlighted moiety).



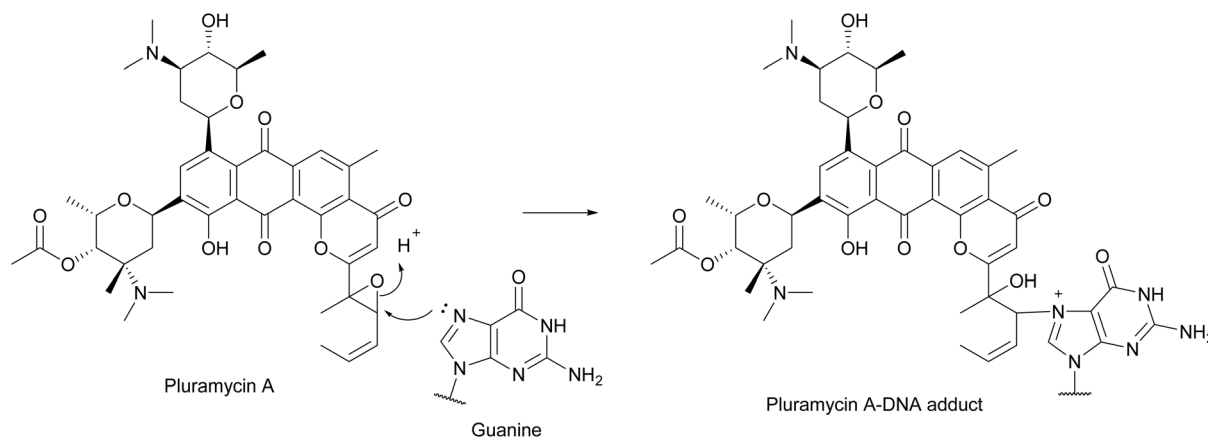


Fig. 33 The DNA alkylation mechanism of pluramycin A.

the nitrosoarea first hydrates and then forms diazene hydroxide, which can act as an electrophile for nucleotide bases in DNA (Fig. 34B).²⁸⁵ Streptozotocin treatment has been reported to methylate at different sites, such as N-7 and O-6 of guanine and N-3 and N-7 of adenine (Fig. 34C).^{286,287} Due to its DNA-alkylating property, streptozotocin presumably induces prophages *via* the SOS pathway discussed above.

4.1.5. Colibactin. Colibactin is a genotoxic metabolite first discovered in 2006, which is synthesized by a 54-kb hybrid nonribosomal peptide synthetase–polyketide synthase (NRPS–PKS) biosynthetic gene cluster (*pks*) in *E. coli*.²⁸⁸ Due to its instability and low yield, colibactin has been recalcitrant to isolation, which precluded efforts to solve its chemical structure.²⁸⁹ Recently, the structure of colibactin has been resolved through a combinatorial approach of genetics, isotope labeling, tandem mass spectrometry, and chemical synthesis (Fig. 35A).^{290,291} Due to its ability to cause DNA double-stranded breaks,²⁸⁸ colibactin has been found to induce the lytic cycle of prophages in a wide range of hosts, such as *pks*[−] *E. coli*, *Salmonella enterica*, *S. aureus*, *Citrobacter rodentium*, and

Enterococcus faecium.²⁹² Colibactin possess a pseudodimeric structure with two reactive cyclopropane warheads located at its two ends (Fig. 35A). These warheads specifically alkylate the N-3 of adenine residues (Fig. 35B) and form inter-strand DNA crosslinks.²⁹³ Since the induction activity is eliminated in a $\Delta recA$ mutant, the prophage induction by colibactin is believed to occur *via* the RecA-dependent SOS pathway.²⁹²

4.1.6. Gilvocarcins. In 1982, gilvocarcins V and M (Fig. 5A, isolated from *Streptomyces arenae* 2064) were found to induce prophages in a biochemical prophage induction assay (BIA).²⁹⁴ In brief, the bacteria used in this assay harbor an engineered λ prophage that produces β -galactosidase as a reporter of prophage induction conditions.¹⁵ However, another study published in the same year found that gilvocarcin V did not induce λ prophage²⁹⁵ in a standard induction assay.¹⁴ This discrepancy was clarified later after the discovery that the DNA-alkylating property of gilvocarcin was light dependent (Fig. 6B).⁶⁰ The previous experiments did not control for light as a variable, explaining the inconsistent results. This photo-activated DNA-alkylating activity of gilvocarcin has been shown to cause DNA

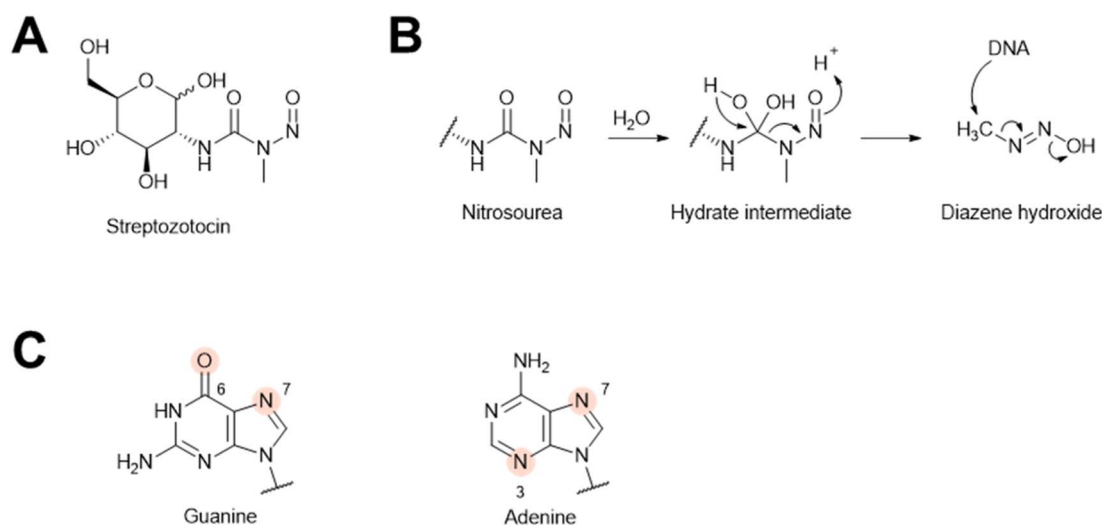


Fig. 34 The prophase induction mechanism of streptozotocin. (A) Chemical structure of streptozotocin. (B) DNA alkylation mechanism of the nitrosoarea functional group. (C) Possible methylation sites on purines by streptozotocin are highlighted.



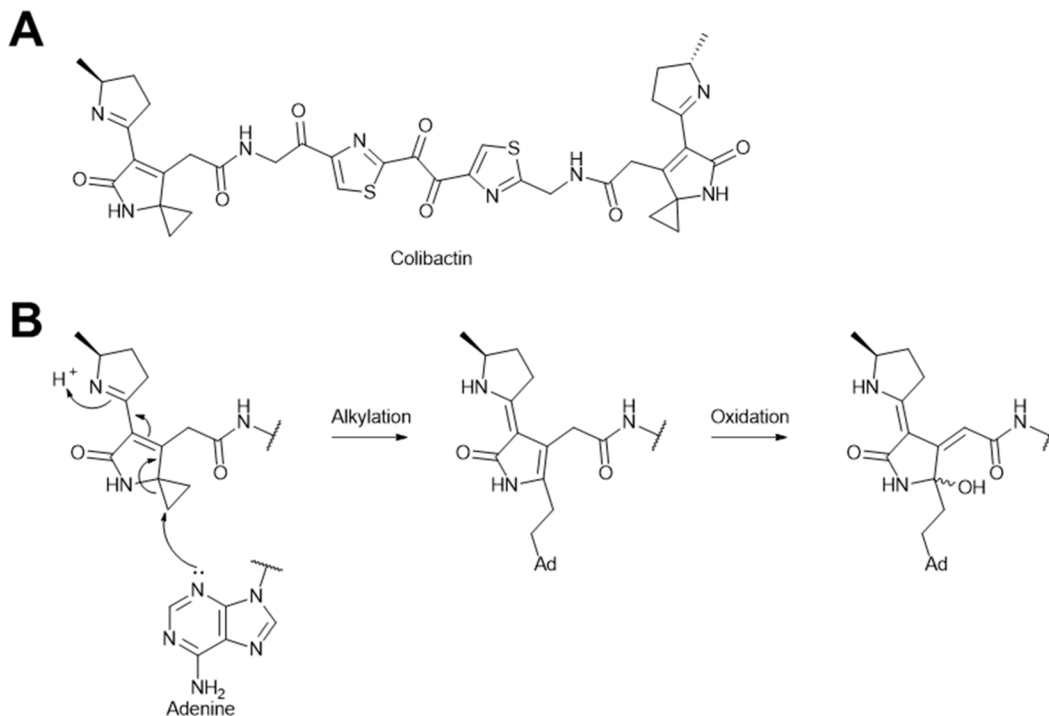


Fig. 35 The prophage induction mechanism of colibactin. (A) Chemical structure of colibactin. (B) DNA-alkylation mechanism of colibactin.

damage both *in vitro* and in cells.^{296–298} This damage likely triggers the SOS response in host bacteria to induce prophages to enter their lytic cycle through similar mechanisms as discussed above.

4.1.7. Bleomycins. Bleomycins are a family of glycopeptide antibiotics with excellent antitumor activities.²⁹⁹ In 1956, phleomycins in this family were first isolated from *Streptomyces*³⁰⁰ as a mixture of 12 structurally related components that only differ at the C-terminus of the peptide backbone (Fig. 36A).³⁰¹ Following the discovery of phleomycins, bleomycins were isolated from *Streptomyces verticillus* as a structurally related mixture with A2 and B2 as the major components.^{302,303} Shortly after their discoveries, both phleomycins and bleomycins were reported to induce λ prophage in *E. coli*.^{14,304} Bleomycins also induced PBSH prophage in *B. subtilis*.³⁰⁴ Other members in this family, such as tallysomycins A and B (Fig. 36) isolated from *Streptomyces*, also induced λ prophage in *E. coli*.³⁰⁵

The prophage induction activity of bleomycins is presumably due to their DNA-damaging mechanisms. The members in the bleomycin family are characterized by a metal-binding domain, a carbohydrate domain, and a DNA-binding domain connected to the former two domains through a linker (Fig. 36A).³⁰⁶ The metal-binding domain can complex with redox-active metal ions to form activated bleomycins, which abstract the 4' hydrogen atom from a deoxyribose residue in DNA, generating DNA strand scission or a 4'-oxidized abasic site (Fig. 36B).^{306–308} These DNA damaging reactions could plausibly trigger the SOS pathway in the bacterial hosts, thus leading to prophage induction through mechanisms discussed earlier.

Beyond prophage induction, bleomycin was also found to inhibit the reproduction of T7 phage on *E. coli*, despite a shorter

latent period.³⁰⁹ The detailed mechanism of such result is still unclear, but it is possibly related to the DNA degradation caused by bleomycin.

4.1.8. Eneidiynes. Eneidyne natural products are anticancer antibiotics with a distinct unsaturated core comprising two acetylenic groups conjugated to a double bond or an incipient double bond.^{299,310} Neocarzinostatin (Fig. 37), the first enediyne antibiotic, was isolated from *Streptomyces carzinostaticus* in 1965 and was reported to induce λ prophage into its lytic cycle.³¹¹ In a search of novel antitumor agents using the BIA experiment in 1989, calicheamicins (Fig. 37) with prophage induction properties were isolated from *Micromonospora echinosporain*.³¹²

Eneidiynes are known to cause DNA damage through a radical-mediated mechanism.^{299,310} For example, the enediyne structures in both neocarzinostatin³¹³ and calicheamicins³¹⁰ can be activated to yield a diradical intermediate (Fig. 37), which abstracts hydrogen atoms from the deoxyribose backbone thus leading to DNA cleavage. DNA damage caused by enediynes likely triggers the SOS response in bacterial hosts, inducing prophages to enter their lytic cycle.

4.1.9. Streptonigrin. Streptonigrin is an aminoquinone antibiotic (Fig. 38A) with antitumor properties that was first isolated from *Streptomyces flocculus* in 1959.³¹⁴ Shortly after its discovery, streptonigrin was reported to induce the lytic cycle in λ and P22 prophages in *E. coli*.^{13,315} The prophage induction activity of streptonigrin is presumably due to its DNA-damaging properties,³¹⁵ which relies on the redox nature of the hydroquinone moiety (Fig. 38B).³¹⁶ The dipyrindyl moiety in streptonigrin can complex with Fe^{2+} , and under aerobic conditions a ferryl radical can be formed (Fig. 38B).³¹⁶ Due to the DNA



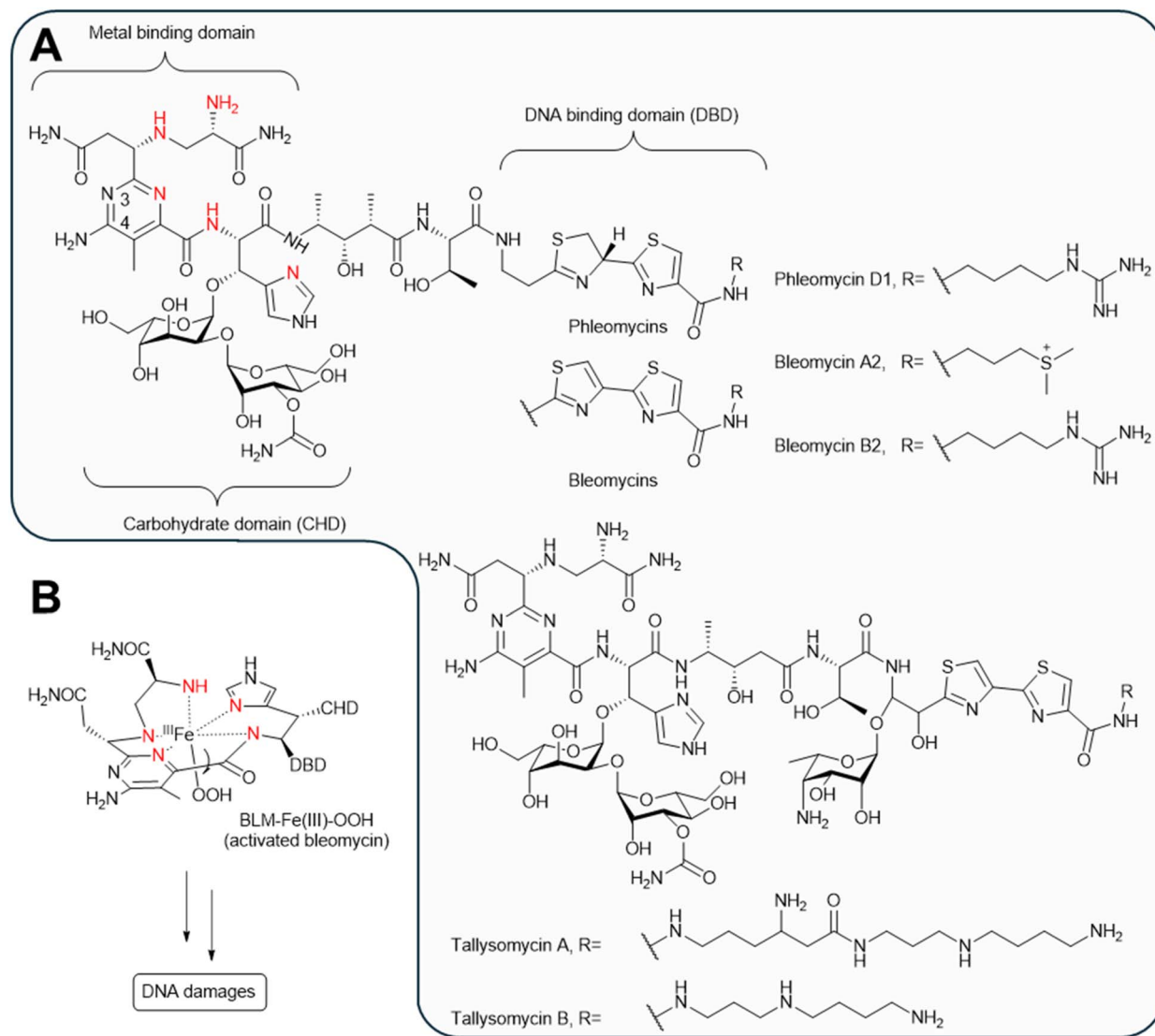


Fig. 36 The prophage induction mechanism of bleomycins. (A) Chemical structures of prophage inducing molecules from the bleomycin family. Metal-coordinating residues are colored red. Phleomycin D1 is shown as a representative phleomycin. Bleomycin A2 and B2 are shown as representative bleomycins. (B) Activated bleomycin is key to the DNA degradation activity.

binding ability of the streptonigrin-Fe complex, the ferryl radical is in proximity with the DNA, inducing DNA damage (Fig. 38B).³¹⁶ Thus, streptonigrin likely induces prophages through an SOS-mediated pathway following the DNA damage, as discussed for the DNA-alkylating agents in the above section.

4.1.10. Xanthomylin. Xanthomylin belongs to the tetracycline class of antibiotics and was first isolated from *Streptomyces* in 1948,³¹⁷ as a tautomeric mixture of its A and B forms (Fig. 39).³¹⁸ In 1964, it was reported that xanthomylin triggered λ prophage induction.¹⁴ However, the exact prophage induction mechanism of xanthomylin is still unclear. It was shown that xanthomylin can cause strand scission in PM2 phage DNA *in vitro*, which is presumably due to the free radicals generated by the quinone moiety.³¹⁹ Therefore, xanthomylin might trigger the lytic cycle through DNA damage-associated SOS-dependent pathways as discussed above.

4.1.11. Griseoluteins. Griseolutein is a phenazine antibiotic that was first isolated from *Streptomyces griseoluteus* in 1950 as a mixture of both A and B forms (Fig. 40).³²⁰ Shortly after its discovery, griseolutein was found to induce λ prophage in *E. coli*.¹⁴ In a later study, a structural analog of griseolutein, pelagiomicin A (Fig. 40),³²¹ was identified from marine bacteria through the BIA experiment.³²² This result suggested that griseolutein and pelagiomicin A might induce prophages into their lytic cycles by generating DNA damage, as the BIA assay specifically detects cellular DNA damage.¹⁵ Both griseolutein and pelagiomicin A feature a phenazine moiety that can cause DNA damage through an iron-dependent pathway.³²³

4.2. Induce redox stress (pyocyanin)

Pyocyanin is a common metabolite produced by *P. aeruginosa* (Fig. 41) with a phenazine core structure.³²⁴ Recently, pyocyanin was shown to induce the lytic cycle in a *S. aureus* prophage.³²⁵



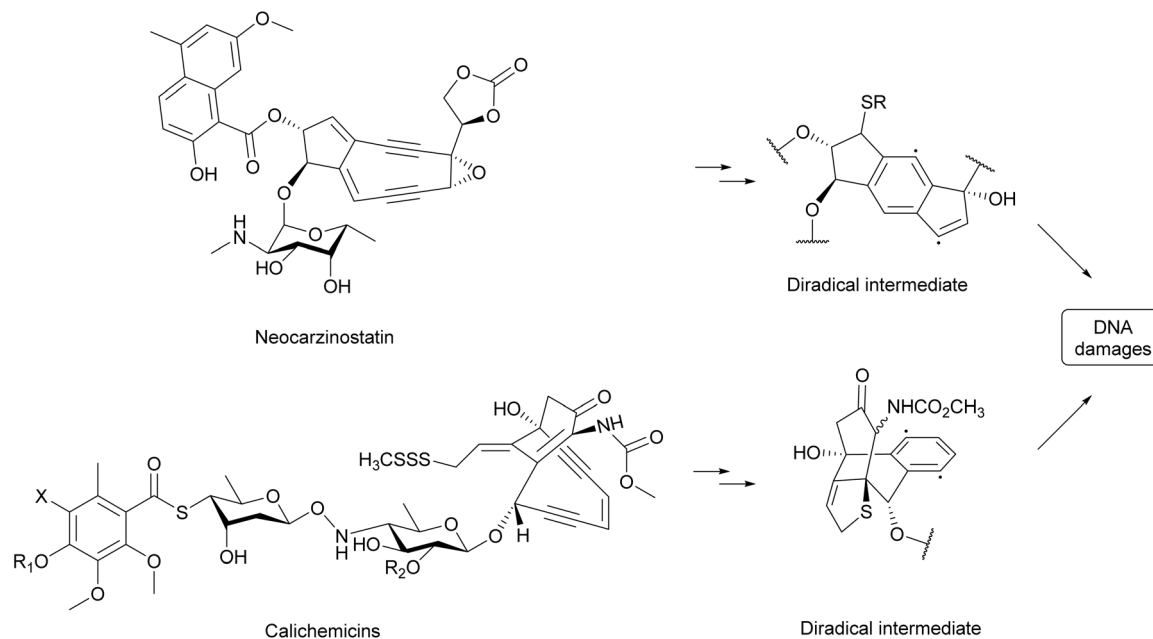


Fig. 37 The prophage induction mechanism of enediynes. Chemical structure of prophage inducing enediynes are shown. The diradical intermediates are essential for the DNA damaging activities of enediynes.

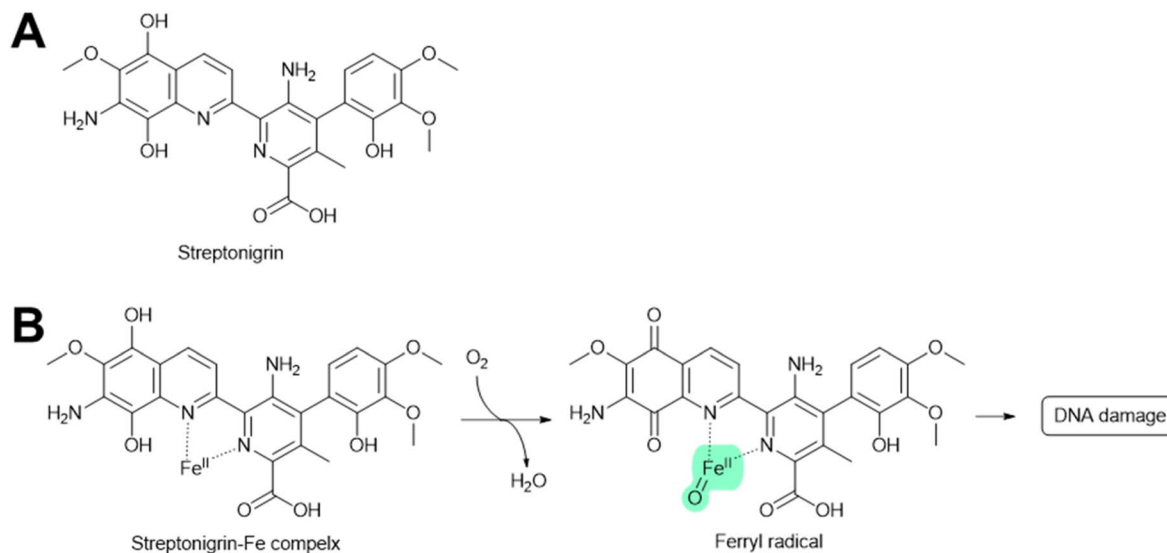


Fig. 38 The prophage induction mechanism of streptonigrin. (A) Chemical structure of streptonigrin. (B) Mechanism of formation of DNA-damaging complex from streptonigrin.

The authors found that pyocyanin induced the prophage through an SOS-independent mechanism,³²⁵ in contrast to the DNA-damaging agents discussed above. Under pyocyanin treatment, multiple oxidative stress response genes in *S. aureus* cells were upregulated, indicating that pyocyanin induced a cellular oxidative stress (Fig. 30 and 41).³²⁵ Notably, the prophage induction by pyocyanin is selective for only certain phages and host strains.³²⁵ In comparison, oxidative stress caused by hydrogen peroxide promiscuously induces many more phages, presumably through oxidative DNA damage. Pyocyanin-induced oxidative stress in the host cells may induce

the lytic cycle through a new mechanism different from the classic de-repression of lytic genes *via* DNA damage.³²⁵ It is surprising that another class of phenazine-containing molecules, griseolutesins (Fig. 40), was shown to cause prophage induction mainly through the DNA damaging pathway as discussed above. Future work could determine which functional groups and/or cellular conditions dictate the different prophage-inducing mechanisms between pyocyanin and griseolutesins. This discovery also implies that a distinct lytic cycle repression mechanism may be encoded by the pyocyanin-sensitive prophages. They may be uniquely de-repressed in an



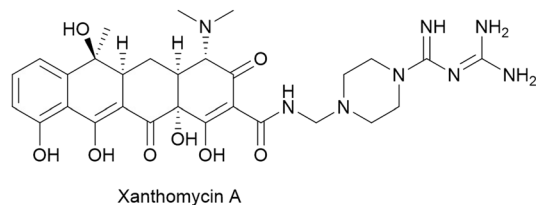


Fig. 39 Chemical structure of xanthomycin A.

oxidative cellular environment. Unraveling a novel de-repression mechanism could advance phage biology and open new avenues for the discovery of prophage-inducing molecules. Since most prophage induction experiments have focused on the λ prophage, there may be many other mechanisms and inducers yet to discover.

4.3. Regulate quorum sensing

Another mechanism evolved by prophages to determine the optimal time to exit the host cell is to sense the density of nearby host cells. It would only be advantageous for a prophage to leave its host cell if there are plenty of uninfected hosts nearby. Therefore, some prophages have evolved the ability to detect host-encoded quorum sensing (QS) signals. In some cases, prophages even encode their own QS signal (Fig. 30) to assess if nearby hosts have already been lysogenized.

4.3.1. Host-encoded QS signals. Group behavior in bacteria is frequently regulated by self-produced QS signal molecules.³²⁶ Since QS signals accumulate as bacterial density increases, a high level of a QS molecule would signal the presence of a high density of hosts for phage infection. Some prophages have leveraged this correlation of QS signal concentration and host density to regulate entry into their lytic cycles. The first example reported was *Pseudomonas* quinolone signal (PQS, Fig. 42A),³²⁷ a QS signal produced by *P. aeruginosa*.³²⁸ PQS was shown to induce prophage entry into its lytic cycle in *Pseudomonas putida*.³²⁷ However, the molecular mechanism underlying the prophage induction by PQS is still elusive.

A second example was autoinducer-2 (AI-2, Fig. 42A).³²⁹ This signal was initially identified as a QS signal in *Vibrio harvey*^{330,331} but was later found to be a widespread QS signal produced by many bacteria.³²⁶ In 2015, it was found that AI-2 can induce multiple prophages in *E. faecalis* (a bacterium that

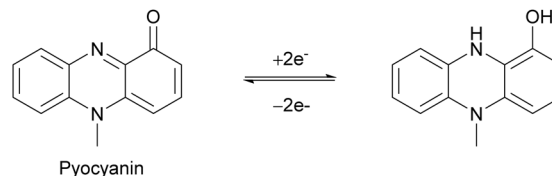


Fig. 41 Redox equilibrium of pyocyanin.

uses the AI-2 QS signal), although the mechanism of action is still unclear.³²⁹

A third example is 3,5-dimethylpyrazin-2-ol (DPO, Fig. 42A),³³² a QS signal in *V. cholerae*.³³³ DPO can induce the *Vibrio parahaemolyticus* VP882 prophage to enter its lytic cycle.³³² VP882 encodes a DPO-binding QS receptor ($VqmA_{\text{phage}}$), which shares homology with the host QS receptor.³³² Upon DPO binding, $VqmA_{\text{phage}}$ induces the expression of an anti-repressor, which was named “quorum-triggered inactivator of cI protein” (Qtip).³³² Qtip then inactivates the lytic gene repressor, cI, thus triggering the phage lytic program.³³² The phage particles then disperse to infect the dense population of nearby bacteria.

4.3.2. Phage-encoded QS signals. In addition to hijacking host-encoded QS signals, some phages also encode their own QS signal to coordinate the lysis-lysogeny decision.³³⁴ This strategy can inform the phage if its nearby host population has already been lysogenized—in which case there is no benefit of trying to “re-infect” those hosts. For example, phages of the SPbeta group encode a six amino-acid-long peptide named “arbitrium” (Fig. 42B), which can regulate their lysis/lysogeny decision.³³⁴ The arbitrium system consists of three genes: *aimP*, encoding the arbitrium peptide; *aimR*, the arbitrium peptide receptor and transcription factor; and *aimX*, which encodes an AimR-regulated non-coding RNA that represses lysogeny.³³⁴ Since *aimP* and *aimR* reside in the same operon, AimP and AimR are simultaneously expressed upon phage infection.³³⁴ AimP is a precursor peptide that is secreted and processed extracellularly into the mature arbitrium peptide. On the other hand, AimR forms a dimer at low phage density, and activates the expression of *aimX*, which represses lysogeny.³³⁴ Arbitrium peptides can accumulate in the medium and be internalized into the host bacteria by an oligopeptide permease transporter.³³⁴ A high density of extracellular peptide (indicating nearby lysogenized cells) will lead to an elevated

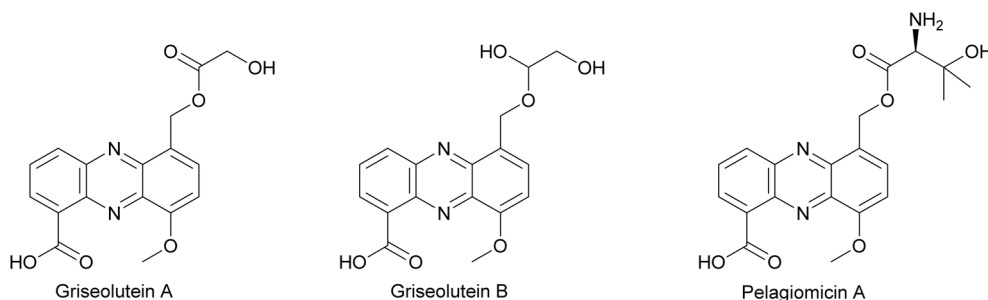


Fig. 40 Chemical structures of griseolutesins.



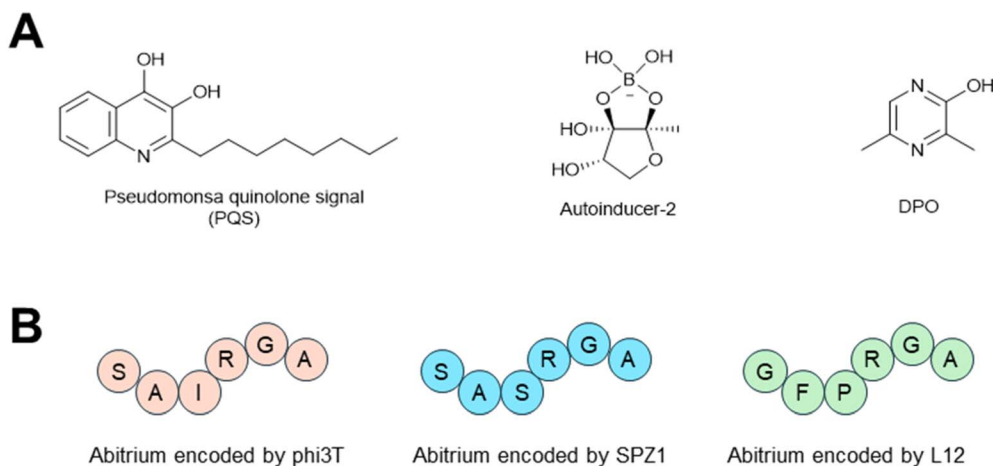


Fig. 42 Chemical structures of prophage-regulating quorum sensing (QS) signals. (A) Host-encoded QS signals that can induce the lytic cycle in prophages. (B) Example arbitrium peptides that signal phages to enter and remain in their temperate prophage states.

intracellular level of arbitrium peptides, which then bind to AimR and antagonize its activation of *aimX* expression, thus biasing phages to enter (and remain in) the lysogenic cycle.³³⁴

5. Conclusions and outlook

As shown through this manuscript, the long history of natural product research has revealed many metabolites that influence phage behavior. However, the ecological and therapeutic implications of the antagonisms and synergies between natural products and phages are still largely unclear. First, it is worth investigating why the genes encoding these phage-modulating compounds are preserved along the evolutionary path. For example, are some bacterial metabolites that are traditionally thought of as antibiotics actually produced to modulate phage predation as their primary role? Second, with respect to phage-based interventions (*e.g.* phage therapy), an expanded knowledge of the phage-interacting “metabolome” in the actual application settings would help to understand and overcome factors that may diminish phage efficacy. On the other hand, future discoveries of phage-promoting natural products may open new avenues as adjuvants to improve phage efficacy.

Despite many early discoveries of natural products that modulate phage activities, technical limitations and a poor understanding of phage biology obfuscated the molecular mechanisms behind the natural product–phage interactions. In some cases, the mechanisms of action can be speculated from the metabolites’ antibiotics or antitumor mechanisms, but generally, elucidation of the phage-influencing mechanisms still requires further investigation.

The recent discovery of natural products inhibiting the Thoiris anti-phage system²⁴⁶ suggests that natural inhibitors against many other anti-phage systems may exist. It is possible that microbes have evolved genes to produce such inhibitors to sensitize their neighbors to phages. This behavior would confer a competitive advantage to the producer, as reported in a recent study.²²⁰ Furthermore, a recent metagenomics study has revealed many biosynthetic gene clusters (BGCs) encoded on

phage genomes.³³⁵ In addition to the proposed functions benefiting the host bacteria,³³⁵ the natural products encoded by these BGCs might also modulate phage activities, which requires further investigation. Therefore, we believe that nature is filled with phage-produced and phage-influencing natural products—many of which are yet-uncovered or incompletely understood.

6. Author contributions

Conceptualization – ZZ, JPG; funding acquisition – JPG; investigation – ZZ, JPG; project administration – JPG; supervision – JPG; visualization – ZZ, JPG; writing (original draft) – ZZ; writing (reviewing & editing) – ZZ, JPG.

7. Conflicts of interest

The authors declare no conflicts of interest.

8. Data availability

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

9. Acknowledgments

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10. References

- 1 G. Ofir and R. Sorek, *Cell*, 2018, **172**, 1260–1270.
- 2 F. d’Herelle, *Bull. N. Y. Acad. Med.*, 1931, **7**, 329–348.
- 3 T. Parfitt, *Lancet*, 2005, **365**, 2166–2167.
- 4 E. C. Keen, *Bioessays*, 2015, **37**, 6–9.
- 5 A. D. Hershey and M. Chase, *J. Gen. Physiol.*, 1952, **36**, 39–56.



- 6 S. Brenner, F. Jacob and M. Meselson, *Nature*, 1961, **190**, 576–581.
- 7 H. O. Smith and K. W. Welcox, *J. Mol. Biol.*, 1970, **51**, 379–391.
- 8 W. Jaroszewicz, J. Morcinek-Orłowska, K. Pierzynowska, L. Gaffke and G. Węgrzyn, *FEMS Microbiol. Rev.*, 2021, **46**, fuab052.
- 9 M. Jinek, K. Chylinski, I. Fonfara, M. Hauer, J. A. Doudna and E. Charpentier, *Science*, 2012, **337**, 816–821.
- 10 J. M. A. Blair, M. A. Webber, A. J. Baylay, D. O. Ogbolu and L. J. V. Piddock, *Nat. Rev. Microbiol.*, 2015, **13**, 42–51.
- 11 R. M. Dedrick, C. A. Guerrero-Bustamante, R. A. Garlena, D. A. Russell, K. Ford, K. Harris, K. C. Gilmour, J. Soothill, D. Jacobs-Sera, R. T. Schooley, G. F. Hatfull and H. Spencer, *Nat. Med.*, 2019, **25**, 730–733.
- 12 M. A. Verini and M. Ghione, *Chemotherapy*, 1964, **9**, 144–157.
- 13 J. Lein, B. Heinemann and A. Gourevitch, *Nature*, 1962, **196**, 783–784.
- 14 K. E. Price, R. E. Buck and J. Lein, *Appl. Microbiol.*, 1964, **12**, 428–435.
- 15 R. K. Elespuru and M. B. Yarmolinsky, *Environ. Mutagen.*, 1979, **1**, 65–78.
- 16 A. M. Comeau, F. Tétart, S. N. Trojet, M.-F. Prère and H. M. Krisch, *PLoS One*, 2007, **2**, e799.
- 17 M. Kim, Y. Jo, Y. J. Hwang, H. W. Hong, S. S. Hong, K. Park and H. Myung, *Appl. Environ. Microbiol.*, 2018, **84**, e02085–02018.
- 18 C. G. Liu, S. I. Green, L. Min, J. R. Clark, K. C. Salazar, A. L. Terwilliger, H. B. Kaplan, B. W. Trautner, R. F. Ramig and A. W. Maresso, *mBio*, 2020, **11**, DOI: [10.1128/mbio.01462-20](https://doi.org/10.1128/mbio.01462-20).
- 19 P. Nicholls, J. R. Clark, C. G. Liu, A. Terwilliger and A. W. Maresso, *Infect. Immun.*, 2023, **91**, e00065–00023.
- 20 J. M. Ribeiro, G. N. Pereira, R. K. Kobayashi and G. Nakazato, *Future Microbiol.*, 2020, **15**, 767–777.
- 21 J. León-Félix and C. Villicaña, *J. Bacteriol.*, 2021, **203**, DOI: [10.1128/jb.00687-00620](https://doi.org/10.1128/jb.00687-00620).
- 22 A. Hardy, L. Kever and J. Frunzke, *Trends Microbiol.*, 2023, **31**, 92–106.
- 23 K. Hatano, E. Higashide, M. Shibata, Y. Kameda, S. Horii and K. Mizuno, *Agric. Biol. Chem.*, 1980, **44**, 1157–1163.
- 24 R. A. Altenbern, *J. Bacteriol.*, 1953, **65**, 288–292.
- 25 S. Kronheim, M. Daniel-Ivad, Z. Duan, S. Hwang, A. I. Wong, I. Mantel, J. R. Nodwell and K. L. Maxwell, *Nature*, 2018, **564**, 283–286.
- 26 H. Shomar, F. Tesson, M. Guillaume, V. Ongenae, M. Le Bot, H. Georjon, E. Mordret, L. Zhang, G. P. van Wezel, D. Rozen, A. Briegel, S. Zirah, D. Claessen, Y. Li and A. Bernheim, *bioRxiv*, 2024, preprint, DOI: [10.1101/2024.06.26.600839](https://doi.org/10.1101/2024.06.26.600839).
- 27 J. J. Champoux, *Annu. Rev. Biochem.*, 2001, **70**, 369–413.
- 28 F. Strelitz, H. Flon, U. Weiss and I. N. Asheshov, *J. Bacteriol.*, 1956, **72**, 90–94.
- 29 B. Parisi and A. Soller, *Giorno. Microbiol.*, 1964, **12**, 183.
- 30 A. Di Marco, M. Gaetani, P. Orezzi, B. M. Scarpinato, R. Silvestrini, M. Soldati, T. Dasdia and L. Valentini, *Nature*, 1964, **201**, 706–707.
- 31 J. Morita, A. Tanaka, T. Komano and T. Oki, *Agric. Biol. Chem.*, 1979, **43**, 2629–2631.
- 32 F. Arcamone, G. Cassinelli, G. Fantini, A. Grein, P. Orezzi, C. Pol and C. Spalla, *Biotechnol. Bioeng.*, 1969, **11**, 1101–1110.
- 33 T. Oki, Y. Matsuzawa, A. Yoshimoto, K. Numata and I. Kitamura, *J. Antibiot.*, 1975, **28**, 830–834.
- 34 W. J. Pigram, W. Fuller and L. D. Hamilton, *Nat. New Biol.*, 1972, **235**, 17–19.
- 35 G. J. Quigley, A. H. Wang, G. Ughetto, G. van der Marel, J. H. van Boom and A. Rich, *Proc. Natl. Acad. Sci. U. S. A.*, 1980, **77**, 7204–7208.
- 36 A. H. J. Wang, G. Ughetto, G. J. Quigley and A. Rich, *Biochemistry*, 1987, **26**, 1152–1163.
- 37 C. A. Frederick, L. D. Williams, G. Ughetto, G. A. Van der Marel, J. H. Van Boom, A. Rich and A. H. J. Wang, *Biochemistry*, 1990, **29**, 2538–2549.
- 38 M. F. Goodman, M. J. Bessman and N. R. Bachur, *Proc. Natl. Acad. Sci. U. S. A.*, 1974, **71**, 1193–1196.
- 39 D. C. Ward, E. Reich and I. H. Goldberg, *Science*, 1965, **149**, 1259–1263.
- 40 J. Marinello, M. Delcuratolo and G. Capranico, *Int. J. Mol. Sci.*, 2018, **19**, 3480.
- 41 J. C. Wang, *Nat. Rev. Mol. Cell Biol.*, 2002, **3**, 430–440.
- 42 L. F. Liu, C.-C. Liu and B. M. Alberts, *Nature*, 1979, **281**, 456–461.
- 43 Y. Xin, R. Xian, Y. Yang, J. Cong, Z. Rao, X. Li and Y. Chen, *Nat. Commun.*, 2024, **15**, 8719.
- 44 K. M. Tewey, T. C. Rowe, L. Yang, B. D. Halligan and L. F. Liu, *Science*, 1984, **226**, 466–468.
- 45 A. Someya and N. Tanaka, *J. Antibiot.*, 1979, **32**, 839–845.
- 46 S. X. Van Ravenstein, K. P. Mehta, T. Kavlashvili, J. A. W. Byl, R. Zhao, N. Osheroff, D. Cortez and J. M. Dewar, *EMBO J.*, 2022, **41**, e110632.
- 47 S. Kondo, T. Wakashiro, M. Hamada, K. Maeda and T. Takeuchi, *J. Antibiot.*, 1970, **23**, 354–359.
- 48 N. Tanaka, *J. Antibiot.*, 1970, **23**, 523–530.
- 49 M. Hansen and L. Hurley, *J. Am. Chem. Soc.*, 1995, **117**, 2421–2429.
- 50 D. Sun, M. Hansen and L. Hurley, *J. Am. Chem. Soc.*, 1995, **117**, 2430–2440.
- 51 P. L. Hamilton and D. P. Arya, *Nat. Prod. Rep.*, 2012, **29**, 134–143.
- 52 C. Fischer, F. Lipata and J. Rohr, *J. Am. Chem. Soc.*, 2003, **125**, 7818–7819.
- 53 F. Strelitz, H. Flon and I. N. Asheshov, *J. Bacteriol.*, 1955, **69**, 280–283.
- 54 K. Shishido, K.-i. Joho, M. Uramoto, K. Isono and T. Jain, *Biochem. Biophys. Res. Commun.*, 1986, **136**, 885–890.
- 55 L. R. McGee and R. Misra, *J. Am. Chem. Soc.*, 1990, **112**, 2386–2389.
- 56 F. Tomita, K. Takahashi and T. Tamaoki, *J. Antibiot.*, 1982, **35**, 1038–1041.



- 57 T. T. Wei, K. M. Byrne, D. Warnick-Pickle and M. Greenstein, *J. Antibiot.*, 1982, **35**, 545–548.
- 58 B. Muralikrishnan, L. K. Edison, A. Dusthacker, G. R. Jijimole, R. Ramachandran, A. Madhavan and R. A. Kumar, *J. Antibiot.*, 2022, **75**, 226–235.
- 59 J. Zhang, P. Liu, J. Chen, D. Yao, Q. Liu, J. Zhang, H.-W. Zhang, E. L.-H. Leung, X.-J. Yao and L. Liu, *Pharmacol. Res.*, 2023, **187**, 106565.
- 60 R. K. Elespuru and S. K. Gonda, *Science*, 1984, **223**, 69–71.
- 61 F. Strelitz, H. Flon and I. N. Asheshov, *Proc. Natl. Acad. Sci. U. S. A.*, 1955, **41**, 620–624.
- 62 D. I. Shiriaev, A. A. Sofronova, E. A. Berdnikovich, D. A. Lukianov, E. S. Komarova, V. I. Marina, Y. V. Zakalyukina, M. V. Biryukov, T. P. Maviza, Y. A. Ivanenkov, P. V. Sergiev, I. A. Osterman and O. A. Dontsova, *Antimicrob. Agents Chemother.*, 2021, **65**, DOI: [10.1128/aac.00777-00720](https://doi.org/10.1128/aac.00777-00720).
- 63 A. C. Finlay, F. A. Hochstein, B. A. Sobin and F. X. Murphy, *J. Am. Chem. Soc.*, 1951, **73**, 341–343.
- 64 F. Arcamone, F. Bizioli, G. Canevazzi and A. Grein, *German Pat.*, DE1039198, 1958.
- 65 A. Di Marco, M. Ghione, A. Sanfilippo and E. Morvillo, *Experientia*, 1963, **19**, 134–136.
- 66 A. Di Marco, A. Migliacci, M. Ghione, E. Morvillo and A. Sanfilippo, *Giorno. Microbiol.*, 1963, **11**, 87.
- 67 F. M. Schabel, W. R. Laster, R. W. Brockman and H. E. Skipper, *Proc. Soc. Exp. Biol. Med.*, 1953, **83**, 1–3.
- 68 K. Watanabe, *J. Antibiot.*, 1956, **9**, 102–107.
- 69 Y. Becker, Y. Asher and Z. Zakay-Rones, *Antimicrob. Agents Chemother.*, 1972, **1**, 483–488.
- 70 M. L. Kopka, C. Yoon, D. Goodsell, P. Pjura and R. E. Dickerson, *Proc. Natl. Acad. Sci. U. S. A.*, 1985, **82**, 1376–1380.
- 71 M. Coll, C. A. Frederick, A. H. Wang and A. Rich, *Proc. Natl. Acad. Sci. U. S. A.*, 1987, **84**, 8385–8389.
- 72 R. M. Wartell, J. E. Larson and R. D. Wells, *J. Biol. Chem.*, 1974, **249**, 6719–6731.
- 73 B. Puschendorf and H. Grunicke, *FEBS Lett.*, 1969, **4**, 355–357.
- 74 P. Chandra, C. Zimmer and H. Thrum, *FEBS Lett.*, 1970, **7**, 90–94.
- 75 B. Puschendorf, H. Becher, D. Böhländt and H. Grunicke, *Eur. J. Biochem.*, 1974, **49**, 531–537.
- 76 H. A. Küpper, W. T. McAllister and E. K. F. Bautz, *Eur. J. Biochem.*, 1973, **38**, 581–586.
- 77 U. H. Mortensen, T. Stevnsner, S. Krogh, K. Olesen, O. Westergaard and B. J. Bonven, *Nucleic Acids Res.*, 1990, **18**, 1983–1989.
- 78 J. M. Woynarowski, M. McHugh, R. D. Sigmund and T. A. Beerman, *Mol. Pharmacol.*, 1989, **35**, 177.
- 79 U. Hollstein, *Chem. Rev.*, 1974, **74**, 625–652.
- 80 J. I. Kawamata and H. Fujita, *J. Antibiot.*, 1960, **13**, 295–297.
- 81 A. Nakata, M. Sekiguchi and J. Kawamata, *Nature*, 1961, **189**, 246–247.
- 82 M. Furukawa, A. Inoue and K. Asano, *J. Antibiot.*, 1968, **21**, 568–570.
- 83 D. Korn, J. J. Protass and L. Leive, *Biochem. Biophys. Res. Commun.*, 1965, **19**, 473–481.
- 84 W. B. Pritikin and H. Reiter, *J. Virol.*, 1969, **3**, 578–585.
- 85 J. Polatnick and R. B. Arlinghaus, *J. Virol.*, 1967, **1**, 1130–1134.
- 86 H. M. Sobell, S. C. Jain, T. D. Sakore and C. E. Nordman, *Nat. New Biol.*, 1971, **231**, 200–205.
- 87 S. Kamitori and F. Takusagawa, *J. Mol. Biol.*, 1992, **225**, 445–456.
- 88 J. Hurwitz, J. J. Furth, M. Malamy and M. Alexander, *Proc. Natl. Acad. Sci. U. S. A.*, 1962, **48**, 1222–1230.
- 89 J. M. Kirk, *Biochim. Biophys. Acta*, 1960, **42**, 167–169.
- 90 H. M. Sobell, *Proc. Natl. Acad. Sci. U. S. A.*, 1985, **82**, 5328–5331.
- 91 A. Nakata, *Biken J.*, 1962, **5**, 29–43.
- 92 M. Sekiguchi and S. Iida, *Proc. Natl. Acad. Sci. U. S. A.*, 1967, **58**, 2315–2320.
- 93 A. Roy and S. Mitra, *Nature*, 1970, **228**, 365–366.
- 94 B. Gerratana, *Med. Res. Rev.*, 2012, **32**, 254–293.
- 95 K. Arima, M. Kosaka, G. Tamura, H. Imanaka and H. Sakai, *J. Antibiot.*, 1972, **25**, 437–444.
- 96 H. Osada, K. Ishinabe, T. Yano, K. Kajikawa and K. Isono, *Agric. Biol. Chem.*, 1990, **54**, 2875–2881.
- 97 T. Takeuchi, T. Miyamoto, M. Ishizuka, H. Naganawa and S. Kondo, *J. Antibiot.*, 1976, **29**, 93–96.
- 98 K. W. Kohn, V. H. Bono and H. E. Kann, *Biochim. Biophys. Acta Nucleic Acids Protein Synth.*, 1968, **155**, 121–129.
- 99 M. S. Puvvada, S. A. Forrow, J. A. Hartley, P. Stephenson, I. Gibson, T. C. Jenkins and D. E. Thurston, *Biochemistry*, 1997, **36**, 2478–2484.
- 100 M. L. Kopka, D. S. Goodsell, I. Baikalov, K. Grzeskowiak, D. Cascio and R. E. Dickerson, *Biochemistry*, 1994, **33**, 13593–13610.
- 101 J. M. Sayer, B. Pinsky, A. Schonbrunn and W. Washtien, *J. Am. Chem. Soc.*, 1974, **96**, 7998–8009.
- 102 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.
- 103 L. M. Repka, J. R. Chekan, S. K. Nair and W. A. van der Donk, *Chem. Rev.*, 2017, **117**, 5457–5520.
- 104 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.
- 105 A. D. P. V. Staden, W. F. V. Zyl, M. Trindade, L. M. T. Dicks and C. Smith, *Appl. Environ. Microbiol.*, 2021, **87**, e00186–00121.
- 106 K. S. Makarova, Y. I. Wolf, S. Snir and E. V. Koonin, *J. Bacteriol.*, 2011, **193**, 6039–6056.
- 107 S. Doron, S. Melamed, G. Ofir, A. Leavitt, A. Lopatina, M. Keren, G. Amitai and R. Sorek, *Science*, 2018, **359**, eaar4120.
- 108 P.-J. Ceysens, J. De Smet, J. Wagemans, N. Akulenko, E. Klimuk, S. Hedge, M. Voet, H. Hendrix, J. Paeshuysse, B. Landuyt, H. Xu, J. Blanchard, K. Severinov and R. Lavigne, *Viruses*, 2020, **12**, 976.
- 109 J. Clardy, M. A. Fischbach and C. R. Currie, *Curr. Biol.*, 2009, **19**, R437–R441.
- 110 D. Walsh and I. Mohr, *Nat. Rev. Microbiol.*, 2011, **9**, 860–875.
- 111 K. M. Krause, A. W. Serio, T. R. Kane and L. E. Connolly, *Cold Spring Harb. Perspect. Med.*, 2016, **6**, a027029.
- 112 Y. Takahashi and M. Igarashi, *J. Antibiot.*, 2018, **71**, 4–14.
- 113 A. Schatz, E. Bugle and S. A. Waksman, *Proc. Soc. Exp. Biol. Med.*, 1944, **55**, 66–69.
- 114 D. Jones, *J. Bacteriol.*, 1945, **50**, 341–348.
- 115 T. D. Brock, *Biochem. Biophys. Res. Commun.*, 1962, **9**, 184–187.
- 116 T. D. Brock and S. O. Wooley, *Science*, 1963, **141**, 1065–1067.
- 117 T. D. Brock, J. Mosser and B. Peacher, *Microbiology*, 1963, **33**, 9–22.
- 118 J. Schindler, *Folia Microbiol.*, 1964, **5**, 269–276.
- 119 W. D. Jones and J. Greenberg, *J. Gen. Virol.*, 1978, **39**, 555–557.
- 120 Z. Jiang, J. Wei, Y. Liang, N. Peng and Y. Li, *Antibiotics*, 2020, **9**, 714.
- 121 M. Kozak and D. Nathans, *J. Mol. Biol.*, 1972, **70**, 41–55.
- 122 P. Zuo, P. Yu and P. J. J. Alvarez, *Appl. Environ. Microbiol.*, 2021, **87**, e00468–00421.
- 123 L. Kever, A. Hardy, T. Luthe, M. Hünnefeld, C. Gätgens, L. Milke, J. Wiechert, J. Wittmann, C. Moraru, J. Marienhagen, J. Frunzke and G. Storz, *mBio*, 2022, **13**, e00783–00722.
- 124 K. Joy Harrison, J. Beavon and E. Griffin, *Lancet*, 1959, **273**, 908–910.
- 125 D. N. Wilson, *Nat. Rev. Microbiol.*, 2014, **12**, 35–48.
- 126 B. S. Schuwirth, J. M. Day, C. W. Hau, G. R. Janssen, A. E. Dahlberg, J. H. D. Cate and A. Vila-Sanjurjo, *Nat. Struct. Mol. Biol.*, 2006, **13**, 879–886.
- 127 A. P. Carter, W. M. Clemons, D. E. Brodersen, R. J. Morgan-Warren, B. T. Wimberly and V. Ramakrishnan, *Nature*, 2000, **407**, 340–348.
- 128 I. Wohlgenuth, R. Garofalo, E. Samatova, A. N. Güneç, C. Lenz, H. Urlaub and M. V. Rodnina, *Nat. Commun.*, 2021, **12**, 1830.
- 129 M. B. Feldman, D. S. Terry, R. B. Altman and S. C. Blanchard, *Nat. Chem. Biol.*, 2010, **6**, 54–62.
- 130 M. A. Borovinskaya, S. Shoji, K. Fredrick and J. H. Cate, *RNA*, 2008, **14**, 1590–1599.
- 131 T. Matt, C. L. Ng, K. Lang, S.-H. Sha, R. Akbergenov, D. Shcherbakov, M. Meyer, S. Duscha, J. Xie, S. R. Dubbaka, D. Perez-Fernandez, A. Vasella, V. Ramakrishnan, J. Schacht and E. C. Böttger, *Proc. Natl. Acad. Sci. U. S. A.*, 2012, **109**, 10984–10989.
- 132 L. Wang, A. Pulk, M. R. Wasserman, M. B. Feldman, R. B. Altman, J. H. D. Cate and S. C. Blanchard, *Nat. Struct. Mol. Biol.*, 2012, **19**, 957–963.
- 133 M. Kopaczynska, M. Lauer, A. Schulz, T. Wang, A. Schaefer and J.-H. Fuhrhop, *Langmuir*, 2004, **20**, 9270–9275.
- 134 C. Weigel and H. Seitz, *FEMS Microbiol. Rev.*, 2006, **30**, 321–381.
- 135 S. Nechaev and K. Severinov, *Annu. Rev. Microbiol.*, 2003, **57**, 301–322.
- 136 H. Yang, Y. Ma, Y. Wang, H. Yang, W. Shen and X. Chen, *Bioengineered*, 2014, **5**, 300–304.
- 137 W. T. McAllister and C. L. Barrett, *J. Virol.*, 1977, **23**, 543–553.
- 138 A. Tabib-Salazar, B. Liu, A. Shadrin, L. Burchell, Z. Wang, Z. Wang, M. G. Goren, I. Yosef, U. Qimron, K. Severinov, S. J. Matthews and S. Wigneshweraraj, *Nucleic Acids Res.*, 2017, **45**, 7697–7707.
- 139 I. Chopra and M. Roberts, *Microbiol. Mol. Biol. Rev.*, 2001, **65**, 232–260.
- 140 B. M. Duggar, *Ann. N. Y. Acad. Sci.*, 1948, **51**, 177–181.
- 141 H. Brainerd, E. H. Lennette, G. Meiklejohn, H. B. Bruyn and W. H. Clark, *J. Clin. Invest.*, 1949, **28**, 992–1005.
- 142 D. E. Brodersen, W. M. Clemons Jr, A. P. Carter, R. J. Morgan-Warren, B. T. Wimberly and V. Ramakrishnan, *Cell*, 2000, **103**, 1143–1154.
- 143 L. Jenner, A. L. Starosta, D. S. Terry, A. Mikolajka, L. Filonava, M. Yusupov, S. C. Blanchard, D. N. Wilson and G. Yusupova, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 3812–3816.
- 144 T. Dimitriu, E. Kurilovich, U. Łapińska, K. Severinov, S. Pagliara, M. D. Szczelkun and E. R. Westra, *Cell Host Microbe*, 2022, **30**, 31–40.
- 145 B. J. Pons, T. Dimitriu, E. R. Westra and S. van Houte, *Proc. Natl. Acad. Sci. U. S. A.*, 2023, **120**, e2216084120.
- 146 S. Y. Stanley, A. L. Borges, K.-H. Chen, D. L. Swaney, N. J. Krogan, J. Bondy-Denomy and A. R. Davidson, *Cell*, 2019, **178**, 1452–1464.
- 147 J. Ehrlich, Q. R. Bartz, R. M. Smith, D. A. Joslyn and P. R. Burkholder, *Science*, 1947, **106**, 417.
- 148 F. M. Bozeman, C. L. Wisseman, H. E. Hopps and J. X. Danauskas, *J. Bacteriol.*, 1954, **67**, 530–536.
- 149 J.-i. Tomizawa and S. Sunakawa, *J. Gen. Physiol.*, 1956, **39**, 553–565.
- 150 L. V. Crawford, *Virology*, 1959, **7**, 359–374.
- 151 R. Thomas, *Virology*, 1959, **9**, 275–289.
- 152 J. M. Erskine, *Appl. Microbiol.*, 1970, **19**, 707–713.
- 153 A. D. Hershey and N. E. Melechen, *Virology*, 1957, **3**, 207–236.



- 154 J. A. Dunkle, L. Xiong, A. S. Mankin and J. H. D. Cate, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 17152–17157.
- 155 D. Bulkley, C. A. Innis, G. Blaha and T. A. Steitz, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 17158–17163.
- 156 J. Marks, K. Kannan, E. J. Roncase, D. Klepacki, A. Kefi, C. Orelle, N. Vázquez-Laslop and A. S. Mankin, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**, 12150–12155.
- 157 P. L. Friend and H. D. Slade, *J. Virol.*, 1967, **1**, 50–56.
- 158 J. R. Christensen, *Virology*, 1957, **4**, 184–185.
- 159 J. Staunton and B. Wilkinson, *Chem. Rev.*, 1997, **97**, 2611–2630.
- 160 G. J. Mc, R. L. Bunch, R. C. Anderson, H. E. Boaz, E. H. Flynn, H. M. Powell and J. W. Smith, *Antibiot. Chemother.*, 1952, **2**, 281–283.
- 161 H. Hirochika, *Mol. Gen. Genet.*, 1980, **179**, 581–588.
- 162 E. Franck and T. S. Crofts, *npj Antimicrob. Resist.*, 2024, **2**, 3.
- 163 S. A. Waksman and H. B. Woodruff, *Proc. Soc. Exp. Biol. Med.*, 1942, **49**, 207–210.
- 164 T. Miyakawa, N. Shimizu and E. Horigome, *Virus*, 1957, **7**, 400–404.
- 165 C. E. Morgan, Y.-S. Kang, A. B. Green, K. P. Smith, M. G. Dowgiallo, B. C. Miller, L. Chiaraviglio, K. A. Truelson, K. E. Zulauf, S. Rodriguez, A. D. Kang, R. Manetsch, E. W. Yu and J. E. Kirby, *PLoS Biol.*, 2023, **21**, e3002091.
- 166 I. Haupt, R. Hübener and H. Thrum, *J. Antibiot.*, 1978, **31**, 1137–1142.
- 167 I. Haupt, J. Jonák, I. Rychlík and H. Thrum, *J. Antibiot.*, 1980, **33**, 636–641.
- 168 A. Parmeggiani and P. Nissen, *FEBS Lett.*, 2006, **580**, 4576–4581.
- 169 S. M. Prezioso, N. E. Brown and J. B. Goldberg, *Mol. Microbiol.*, 2017, **106**, 22–34.
- 170 H. Wolf and H. Zähner, *Arch. Mikrobiol.*, 1972, **83**, 147–154.
- 171 V. P. Gullo, S. B. Zimmerman, R. S. Dewey, O. Hensens, P. J. Cassidy, R. Oiwa and S. Omura, *J. Antibiot.*, 1982, **35**, 1705–1707.
- 172 R. S. Dewey, J. E. Flor, S. B. Zimmerman, P. J. Cassidy, S. Omura and R. Oiwa, *US Pat.*, 4262002, 1981.
- 173 H. Kakeya, M. Morishita, A. Ikeno, K. Kobinata, T. Yano and H. Osada, *J. Antibiot.*, 1998, **51**, 963–966.
- 174 J. C. Morse, D. Girodat, B. J. Burnett, M. Holm, R. B. Altman, K. Y. Sanbonmatsu, H.-J. Wieden and S. C. Blanchard, *Proc. Natl. Acad. Sci. U. S. A.*, 2020, **117**, 3610–3620.
- 175 L. Vogeley, G. J. Palm, J. R. Mesters and R. Hilgenfeld, *J. Biol. Chem.*, 2001, **276**, 17149–17155.
- 176 H. Brötz-Oesterhelt, D. Beyer, H.-P. Kroll, R. Endermann, C. Ladel, W. Schroeder, B. Hinzen, S. Raddatz, H. Paulsen, K. Henninger, J. E. Bandow, H.-G. Sahl and H. Labischinski, *Nat. Med.*, 2005, **11**, 1082–1087.
- 177 M. Arvanitis, G. Li, D.-D. Li, D. Cotnoir, L. Ganley-Leal, D. W. Carney, J. K. Sello and E. Mylonakis, *PLoS One*, 2016, **11**, e0153912.
- 178 S. Z. Z. Cobongela, M. M. Makatini, P. S. Mdluli and N. R. S. Sibuyi, *Pharmaceutics*, 2022, **14**, 1956.
- 179 K. H. Michel and R. E. Kastner, *US Pat.*, 4492650, 1985.
- 180 H. Osada, T. Yano, H. Koshino and K. Isono, *J. Antibiot.*, 1991, **44**, 1463–1466.
- 181 A. O. Olivares, T. A. Baker and R. T. Sauer, *Nat. Rev. Microbiol.*, 2016, **14**, 33–44.
- 182 R. M. Raju, A. L. Goldberg and E. J. Rubin, *Nat. Rev. Drug Discovery*, 2012, **11**, 777–789.
- 183 M. F. Mabanglo and W. A. Houry, *J. Biol. Chem.*, 2022, **298**, 101781.
- 184 B.-G. Lee, E. Y. Park, K.-E. Lee, H. Jeon, K. H. Sung, H. Paulsen, H. Rübnsamen-Schaeff, H. Brötz-Oesterhelt and H. K. Song, *Nat. Struct. Mol. Biol.*, 2010, **17**, 471–478.
- 185 N. Mulvenna, I. Hantke, L. Burchell, S. Nicod, D. Bell, K. Turgay and S. Wigneshweraraj, *J. Biol. Chem.*, 2019, **294**, 17501–17511.
- 186 R. C. Hider and X. Kong, *Nat. Prod. Rep.*, 2010, **27**, 637–657.
- 187 J. Kramer, Ö. Özkaya and R. Kümmerli, *Nat. Rev. Microbiol.*, 2020, **18**, 152–163.
- 188 Z. Zang, K. J. Park and J. P. Gerdt, *ACS Chem. Biol.*, 2022, **17**, 2396–2403.
- 189 E. E. Wyckoff, B. E. Allred, K. N. Raymond and S. M. Payne, *J. Bacteriol.*, 2015, **197**, 2840–2849.
- 190 B. L. Greene, G. Kang, C. Cui, M. Bennati, D. G. Nocera, C. L. Drennan and J. Stubbe, *Annu. Rev. Biochem.*, 2020, **89**, 45–75.
- 191 C. S. Chiu, S. M. Cox and G. R. Greenberg, *J. Biol. Chem.*, 1980, **255**, 2747–2751.
- 192 Y.-C. Yeh and I. Tessman, *Virology*, 1972, **47**, 767–772.
- 193 K. D. Seed, K. L. Bodi, A. M. Kropinski, H.-W. Ackermann, S. B. Calderwood, F. Qadri, A. Camilli and C. Fraser-Liggett, *mBio*, 2011, **2**, e00334–00310.
- 194 K. Gaur, S. C. Pérez Otero, J. A. Benjamín-Rivera, I. Rodríguez, S. A. Loza-Rosas, A. M. Vázquez Salgado, E. A. Akam, L. Hernández-Matías, R. K. Sharma, N. Alicea, M. Kowaleff, A. V. Washington, A. V. Astashkin, E. Tomat and A. D. Tinoco, *JACS Au*, 2021, **1**, 865–878.
- 195 M. Thoendel, J. S. Kavanaugh, C. E. Flack and A. R. Horswill, *Chem. Rev.*, 2011, **111**, 117–151.
- 196 R. P. Novick and E. Geisinger, *Annu. Rev. Genet.*, 2008, **42**, 541–564.
- 197 B. Wang and T. W. Muir, *Cell Chem. Biol.*, 2016, **23**, 214–224.
- 198 J. Yang, J. Z. Bowring, J. Krusche, E. Lehmann, B. S. Bejder, S. F. Silva, M. S. Bojer, T. Grunert, A. Peschel and H. Ingmer, *Cell Rep.*, 2023, **42**, 113154.
- 199 G. Xia, L. Maier, P. Sanchez-Carballo, M. Li, M. Otto, O. Holst and A. Peschel, *J. Biol. Chem.*, 2010, **285**, 13405–13415.
- 200 S. L. Collins, J. G. Stine, J. E. Bisanz, C. D. Okafor and A. D. Patterson, *Nat. Rev. Microbiol.*, 2023, **21**, 236–247.
- 201 Z. Netter, D. T. Dunham and K. D. Seed, *mBio*, 2023, **14**, e01985–01923.
- 202 K. D. Seed, S. M. Faruque, J. J. Mekalanos, S. B. Calderwood, F. Qadri and A. Camilli, *PLoS Pathog.*, 2012, **8**, e1002917.
- 203 M. Yen, L. S. Cairns and A. Camilli, *Nat. Commun.*, 2017, **8**, 14187.
- 204 H.-B. Li and F. Chen, *J. Chromatogr.*, 2005, **1074**, 107–110.



- 205 J. M. Broniewski, M. A. W. Chisnall, N. M. Høyland-Kroghsbo, A. Buckling and E. R. Westra, *ISME J.*, 2021, **15**, 2465–2473.
- 206 J. M. Budzik, W. A. Rosche, A. Rietsch and G. A. O'Toole, *J. Bacteriol.*, 2004, **186**, 3270–3273.
- 207 K. C. Cady, J. Bondy-Denomy, G. E. Heussler, A. R. Davidson and G. A. O'Toole, *J. Bacteriol.*, 2012, **194**, 5728–5738.
- 208 A. Glessner, R. S. Smith, B. H. Iglewski and J. B. Robinson, *J. Bacteriol.*, 1999, **181**, 1623–1629.
- 209 H. Georjon and A. Bernheim, *Nat. Rev. Microbiol.*, 2023, **21**, 686–700.
- 210 S. J. Hobbs and P. J. Kranzusch, *Annu. Rev. Microbiol.*, 2024, **78**, 255–276.
- 211 D. Cohen, S. Melamed, A. Millman, G. Shulman, Y. Oppenheimer-Shaanan, A. Kacen, S. Doron, G. Amitai and R. Sorek, *Nature*, 2019, **574**, 691–695.
- 212 A. Millman, S. Melamed, G. Amitai and R. Sorek, *Nat. Microbiol.*, 2020, **5**, 1608–1615.
- 213 G. Ofir, E. Herbst, M. Baroz, D. Cohen, A. Millman, S. Doron, N. Tal, D. B. A. Malheiro, S. Malitsky, G. Amitai and R. Sorek, *Nature*, 2021, **600**, 116–120.
- 214 D. Sabonis, C. Avraham, R. B. Chang, A. Lu, E. Herbst, A. Silanskas, D. Vilutis, A. Leavitt, E. Yirmiya, H. C. Toyoda, A. Ruksenaite, M. Zaremba, I. Osterman, G. Amitai, P. J. Kranzusch, R. Sorek and G. Tamulaitiene, *Nature*, 2025, **642**, 467–473.
- 215 F. Rousset, I. Osterman, T. Scherf, A. H. Falkovich, A. Leavitt, G. Amitai, S. Shir, S. Malitsky, M. Itkin, A. Savidor and R. Sorek, *Science*, 2025, **387**, 510–516.
- 216 M. Kazlauskienė, G. Kostiuik, Č. Venclovas, G. Tamulaitis and V. Siksnys, *Science*, 2017, **357**, 605–609.
- 217 O. Niewoehner, C. Garcia-Doval, J. T. Rostøl, C. Berk, F. Schwede, L. Bigler, J. Hall, L. A. Marraffini and M. Jinek, *Nature*, 2017, **548**, 543–548.
- 218 G. Stella and L. Marraffini, *Trends Biochem. Sci.*, 2024, **49**, 28–37.
- 219 N. Tal, B. R. Morehouse, A. Millman, A. Stokar-Avihail, C. Avraham, T. Fedorenko, E. Yirmiya, E. Herbst, A. Brandis, T. Mehlman, Y. Oppenheimer-Shaanan, A. F. A. Keszei, S. Shao, G. Amitai, P. J. Kranzusch and R. Sorek, *Cell*, 2021, **184**, 5728–5739.
- 220 Z. Zang, C. Zhang, K. J. Park, D. A. Schwartz, R. Podicheti, J. T. Lennon and J. P. Gerdt, *Nat. Microbiol.*, 2025, **10**, 362–373.
- 221 *WHO Report on Surveillance of Antibiotic Consumption: 2016–2018 Early Implementation*, World Health Organization, Geneva, 2018.
- 222 A. Fleming, *Br. J. Exp. Pathol.*, 1929, **10**, 226.
- 223 J. Houbraken, J. C. Frisvad and R. A. Samson, *IMA Fungus*, 2011, **2**, 87–95.
- 224 A. P. Krueger, T. Cohn and N. Noble, *Proc. Soc. Exp. Biol. Med.*, 1947, **66**, 204–205.
- 225 K. Tahlan and S. E. Jensen, *J. Antibiot.*, 2013, **66**, 401–410.
- 226 E. P. Abraham, *Rev. Infect. Dis.*, 1979, **1**, 99–105.
- 227 K. M. Papp-Wallace, A. Endimiani, M. A. Taracila and R. A. Bonomo, *Antimicrob. Agents Chemother.*, 2011, **55**, 4943–4960.
- 228 R. B. Sykes and D. P. Bonner, *Rev. Infect. Dis.*, 1985, **7**, S579–S593.
- 229 P. Manohar, M. Madurantakam Royam, B. Loh, B. Bozdogan, R. Nachimuthu and S. Leptihn, *ACS Infect. Dis.*, 2022, **8**, 59–65.
- 230 H. Hadas, M. Einav, I. Fishov and A. Zaritsky, *Microbiology*, 1997, **143**, 179–185.
- 231 M. Pazos and K. Peters, in *Bacterial Cell Walls and Membranes*, ed. A. Kuhn, Springer International Publishing, Cham, 2019, pp. 127–168, DOI: [10.1007/978-3-030-18768-2_5](https://doi.org/10.1007/978-3-030-18768-2_5).
- 232 M. Mora-Ochomogo and C. T. Lohans, *RSC Med. Chem.*, 2021, **12**, 1623–1639.
- 233 D. J. Tipper and J. L. Strominger, *Proc. Natl. Acad. Sci. U. S. A.*, 1965, **54**, 1133–1141.
- 234 D. Lim and N. C. J. Strynadka, *Nat. Struct. Biol.*, 2002, **9**, 870–876.
- 235 Z. Lu, H. Wang, A. Zhang, X. Liu, W. Zhou, C. Yang, L. Guddat, H. Yang, C. J. Schofield and Z. Rao, *Mol. Pharmacol.*, 2020, **97**, 287–294.
- 236 J. Buijs, A. S. M. Dofferhoff, J. W. Mouton, J. H. T. Wagenvoort and J. W. M. van der Meer, *Clin. Microbiol. Infect.*, 2008, **14**, 344–349.
- 237 E. S. C. Rittershaus, S.-H. Baek and C. M. Sassetti, *Cell Host Microbe*, 2013, **13**, 643–651.
- 238 D. A. Schwartz, B. K. Lehmkuhl and J. T. Lennon, *mSphere*, 2022, **7**, e00297–00222.
- 239 M. Łoś, P. Golec, J. M. Łoś, A. Węglewska-Jurkiewicz, A. Czyż, A. Węgrzyn, G. Węgrzyn and P. Neubauer, *BMC Biotechnol.*, 2007, **7**, 13.
- 240 D. Bryan, A. El-Shibiny, Z. Hobbs, J. Porter and E. M. Kutter, *Front. Microbiol.*, 2016, **7**, 1391.
- 241 Z. E. V. Phillips and M. A. Strauch, *CMLS Cell. Mol. Life Sci.*, 2002, **59**, 392–402.
- 242 V. Molle, M. Fujita, S. T. Jensen, P. Eichenberger, J. E. González-Pastor, J. S. Liu and R. Losick, *Mol. Microbiol.*, 2003, **50**, 1683–1701.
- 243 Q. Xiong, H. Zhang, X. Shu, X. Sun, H. Feng, Z. Xu, Á. T. Kovács, R. Zhang and Y. Liu, *npj Biofilms Microbiomes*, 2024, **10**, 117.
- 244 C.-W. Chiu, P.-J. Tsai, C.-C. Lee, W.-C. Ko and Y.-P. Hung, *J. Microbiol. Immunol. Infect.*, 2021, **54**, 1011–1017.
- 245 K. Khanna, J. Lopez-Garrido and K. Pogliano, *Annu. Rev. Microbiol.*, 2020, **74**, 361–386.
- 246 Z. Zang, O. K. Duncan, D. Sabonis, Y. Shi, G. Miraj, I. Fedorova, S. Le, J. Deng, Y. Zhu, Y. Cai, C. Zhang, G. Arya, B. A. Duerkop, H. Liang, J. Bondy-Denomy, T. Ve, G. Tamulaitiene and J. P. Gerdt, *bioRxiv*, 2025, preprint, DOI: [10.1101/2025.02.20.638879](https://doi.org/10.1101/2025.02.20.638879).
- 247 D.-G. Wang, L. Niu, Z.-M. Lin, J.-J. Wang, D.-F. Gao, H.-Y. Sui, Y.-Z. Li and C. Wu, *J. Nat. Prod.*, 2021, **84**, 2744–2748.
- 248 N. A. Frank, M. Széles, S. H. Akone, S. Rasheed, S. Hüttel, S. Frewert, M. M. Hamed, J. Herrmann, S. M. M. Schuler, A. K. H. Hirsch and R. Müller, *Molecules*, 2021, **26**, 4929.
- 249 Y. Wang, J. Zheng, P. Liu, W. Wang and W. Zhu, *Mar. Drugs*, 2011, **9**, 1368–1378.



- 250 S. Chen, J. Wang, Z. Wang, X. Lin, B. Zhao, K. Kaliaperumal, X. Liao, Z. Tu, J. Li, S. Xu and Y. Liu, *Fitoterapia*, 2017, **117**, 71–78.
- 251 P. Samuel, K. T. Ayoob, B. A. Magnuson, U. Wölwer-Rieck, P. B. Jeppesen, P. J. Rogers, I. Rowland and R. Mathews, *J. Nutr.*, 2018, **148**, 1186S–1205S.
- 252 L. Marongiu, E. Brzozowska, J. Brykała, M. Burkard, H. Schmidt, B. Szermer-Olearnik and S. Venturelli, *Sci. Rep.*, 2025, **15**, 1337.
- 253 L. Barksdale and S. B. Arden, *Annu. Rev. Microbiol.*, 1974, **28**, 265–300.
- 254 J. Łoś, S. Zielińska, A. Krajewska, Z. Michalina, A. Małachowska, K. Kwaśnicka and M. Łoś, in *Bacteriophages: Biology, Technology, Therapy*, ed. D. R. Harper, S. T. Abedon, B. H. Burrowes and M. L. McConville, Springer International Publishing, Cham, 2020, pp. 1–33, DOI: [10.1007/978-3-319-40598-8_3-1](https://doi.org/10.1007/978-3-319-40598-8_3-1).
- 255 M. Bednarz, J. A. Halliday, C. Herman and I. Golding, *PLoS One*, 2014, **9**, e100876.
- 256 J. W. Little, in *Regulation of Gene Expression in Escherichia coli*, Springer US, Boston, MA, 1996, pp. 453–479, DOI: [10.1007/978-1-4684-8601-8_22](https://doi.org/10.1007/978-1-4684-8601-8_22).
- 257 K. H. Masłowska, K. Makiela-Dzubska and I. J. Fijalkowska, *Environ. Mol. Mutagen.*, 2019, **60**, 368–384.
- 258 P. P. Khil and R. D. Camerini-Otero, *Mol. Microbiol.*, 2002, **44**, 89–105.
- 259 J. W. Little, *Proc. Natl. Acad. Sci. U. S. A.*, 1984, **81**, 1375–1379.
- 260 M. Kim, H. J. Kim, S. H. Son, H. J. Yoon, Y. Lim, J. W. Lee, Y.-J. Seok, K. S. Jin, Y. G. Yu, S. K. Kim, S. Ryu and H. H. Lee, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**, E2480–E2488.
- 261 R. W. Franck, in *Fortschritte der Chemie organischer Naturstoffe/Progress in the Chemistry of Organic Natural Products*, ed. W. Herz, H. Grisebach and G. W. Kirby, Springer Vienna, Vienna, 1979, pp. 1–45, DOI: [10.1007/978-3-7091-8548-3_1](https://doi.org/10.1007/978-3-7091-8548-3_1).
- 262 S. Wakaki, H. Marumo, K. Tomioka, G. Shimizu, E. Kato, H. Kamada, S. Kudo and Y. Fujimoto, *Antibiot. Chemother.*, 1958, **8**, 228–240.
- 263 N. Otsuji, M. Sekiguchi, T. Iijima and Y. Takagi, *Nature*, 1959, **184**, 1079–1080.
- 264 C. DeBoer, A. Dietz, N. Lummis and G. Savage, *Antimicrob. Agents Annu.*, 1960, 17–22.
- 265 V. N. Iyer and W. Szybalski, *Science*, 1964, **145**, 55–58.
- 266 P. D. Bass, D. A. Gubler, T. C. Judd and R. M. Williams, *Chem. Rev.*, 2013, **113**, 6816–6863.
- 267 M. Sastry, R. Fiala, R. Lipman, M. Tomasz and D. J. Patel, *J. Mol. Biol.*, 1995, **247**, 338–359.
- 268 Y. Wei, A. C. Vollmer and R. A. LaRossa, *J. Bacteriol.*, 2001, **183**, 2259–2264.
- 269 C. C. Nawrat and C. J. Moody, *Nat. Prod. Rep.*, 2011, **28**, 1426–1444.
- 270 S. A. Fusari, R. P. Frohardt, A. Ryder, T. H. Haskell, D. W. Johannessen, C. C. Elder and Q. R. Bartz, *J. Am. Chem. Soc.*, 1954, **76**, 2878–2881.
- 271 Q. R. Bartz, C. C. Elder, R. P. Frohardt, S. A. Fusari, T. H. Haskell, D. W. Johannessen and A. Ryder, *Nature*, 1954, **173**, 72–73.
- 272 J. S. Gots, T. J. Bird and S. Mudd, *Biochim. Biophys. Acta*, 1955, **17**, 449–450.
- 273 W. Kirmse, *Angew. Chem. Int. Ed. Engl.*, 1976, **15**, 251–261.
- 274 D. P. Arya, in *Heterocyclic Antitumor Antibiotics*, ed. M. Lee, Springer Berlin Heidelberg, Berlin, Heidelberg, 2006, pp. 129–152, DOI: [10.1007/7081_018](https://doi.org/10.1007/7081_018).
- 275 M. O'Driscoll, P. Macpherson, Y.-Z. Xu and P. Karran, *Carcinogenesis*, 1999, **20**, 1855–1862.
- 276 S. M. Geisen, C. M. N. Aloisi, S. M. Huber, E. S. Sandell, N. A. Escher and S. J. Sturla, *Chem. Res. Toxicol.*, 2021, **34**, 1518–1529.
- 277 H. E. Kubitschek and R. J. Sepanski, *Mutat. Res. Fund Mol. Mech. Mutagen*, 1982, **94**, 31–38.
- 278 D. M. Williams-Hill, J. Olesen, C. Zucker and H. E. Kubitschek, *Mutat. Res. Fund Mol. Mech. Mutagen*, 1984, **129**, 153–164.
- 279 K. Maeda, T. Takeuchi, K. Nitta, K. Yagishita, R. Utahara, T. Ōsato, M. Ueda, S. Kondō, Y. Okami and H. Umezawa, *J. Antibiot.*, 1956, **9**, 75–81.
- 280 B. Heinemann and A. J. Howard, *Appl. Microbiol.*, 1964, **12**, 234–239.
- 281 D. Sun, M. Hansen, J. J. Clement and L. H. Hurley, *Biochemistry*, 1993, **32**, 8068–8074.
- 282 L. C. Tu, T. Melendy and T. A. Beerman, *Mol. Cancer Ther.*, 2004, **3**, 577–586.
- 283 J. J. Vavra, C. Deboer, A. Dietz, L. J. Hanka and W. T. Sokolski, *Antibiot. Annu.*, 1959, **7**, 230–235.
- 284 R. L. Wurdeman, K. M. Church and B. Gold, *J. Am. Chem. Soc.*, 1989, **111**, 6408–6412.
- 285 C. Avendaño and J. C. Menéndez, in *Medicinal Chemistry of Anticancer Drugs*, ed. C. Avendaño and J. C. Menéndez, Elsevier, Boston, 3rd edn, 2023, pp. 237–290, DOI: [10.1016/B978-0-12-818549-0.00004-2](https://doi.org/10.1016/B978-0-12-818549-0.00004-2).
- 286 K. S. Gates, in *Comprehensive Natural Products Chemistry*, ed. S. D. Barton, K. Nakanishi and O. Meth-Cohn, Pergamon, Oxford, 1999, pp. 491–552, DOI: [10.1016/B978-0-08-091283-7.00074-6](https://doi.org/10.1016/B978-0-08-091283-7.00074-6).
- 287 A. D. Bolzán and M. S. Bianchi, *Mutat. Res. Rev. Mutat. Res.*, 2002, **512**, 121–134.
- 288 J.-P. Nougayrède, S. Homburg, F. Taieb, M. Boury, E. Brzuszkiewicz, G. Gottschalk, C. Buchrieser, J. Hacker, U. Dobrindt and E. Oswald, *Science*, 2006, **313**, 848–851.
- 289 J.-W. Tang, X. Liu, W. Ye, Z.-R. Li and P.-Y. Qian, *Nat. Prod. Rep.*, 2022, **39**, 991–1014.
- 290 M. Xue, C. S. Kim, A. R. Healy, K. M. Wernke, Z. Wang, M. C. Frischling, E. E. Shine, W. Wang, S. B. Herzon and J. M. Crawford, *Science*, 2019, **365**, eaax2685.
- 291 Y. Jiang, A. Stornetta, P. W. Villalta, M. R. Wilson, P. D. Boudreau, L. Zha, S. Balbo and E. P. Balskus, *J. Am. Chem. Soc.*, 2019, **141**, 11489–11496.
- 292 J. E. Silpe, J. W. H. Wong, S. V. Owen, M. Baym and E. P. Balskus, *Nature*, 2022, **603**, 315–320.
- 293 M. R. Wilson, Y. Jiang, P. W. Villalta, A. Stornetta, P. D. Boudreau, A. Carrá, C. A. Brennan, E. Chun, L. Ngo,



- L. D. Samson, B. P. Engelward, W. S. Garrett, S. Balbo and E. P. Balskus, *Science*, 2019, **363**, eaar7785.
- 294 T. T. Wei, J. A. Chan, P. P. Roller, U. Weiss, R. M. Stroshane, R. J. White and K. M. Byrne, *J. Antibiot.*, 1982, **35**, 529–532.
- 295 D. M. Balitz, F. A. O'Herron, J. Bush, D. M. Vyas, D. E. Nettleton, R. E. Grulich, W. T. Bradner, T. W. Doyle, E. Arnold and J. Clardy, *J. Antibiot.*, 1981, **34**, 1544–1555.
- 296 F. P. Gasparro, R. M. Knobler and R. L. Edelson, *Chem.-Biol. Interact.*, 1988, **67**, 255–265.
- 297 Y.-C. Tse-Dinh and L. R. McGee, *Biochem. Biophys. Res. Commun.*, 1987, **143**, 808–812.
- 298 M. J. Peak, J. G. Peak, C. M. Blaumueller and R. K. Elespuru, *Chem.-Biol. Interact.*, 1988, **67**, 267–274.
- 299 U. Galm, M. H. Hager, S. G. Van Lanen, J. Ju, J. S. Thorson and B. Shen, *Chem. Rev.*, 2005, **105**, 739–758.
- 300 K. Maeda, H. Kosaka, K. Yagishita and H. Umezawa, *J. Antibiot.*, 1956, **9**, 82–85.
- 301 T. Ikekawa, F. Iwami, H. Hiranaka and H. Umezawa, *J. Antibiot.*, 1964, **17**, 194–199.
- 302 H. Umezawa, K. Maeda, T. Takeuchi and Y. Okami, *J. Antibiot.*, 1966, **19**, 200–209.
- 303 A. Fujii, T. Takita, K. Maeda and H. Umezawa, *J. Antibiot.*, 1973, **26**, 396–397.
- 304 C. W. Haidle, K. K. Weiss and M. L. Mace, *Biochem. Biophys. Res. Commun.*, 1972, **48**, 1179–1184.
- 305 H. Kawaguchi, H. Tsukiura, K. Tomita, M. Konishi and K. Saito, *J. Antibiot.*, 1977, **30**, 779–788.
- 306 A. D. Steele, E. Kalkreuter, G. Pan, S. Meng and B. Shen, in *Comprehensive Natural Products III*, ed. H.-W. Liu and T. P. Begley, Elsevier, Oxford, 2020, pp. 284–335, DOI: [10.1016/B978-0-12-409547-2.14669-4](https://doi.org/10.1016/B978-0-12-409547-2.14669-4).
- 307 J. Chen and J. Stubbe, *Nat. Rev. Cancer*, 2005, **5**, 102–112.
- 308 S. M. Hecht, *J. Nat. Prod.*, 2000, **63**, 158–168.
- 309 K. Shishido, A. Watarai, S. Naito and T. Ando, *J. Antibiot.*, 1975, **28**, 676–680.
- 310 K. C. Nicolaou, A. L. Smith and E. W. Yue, *Proc. Natl. Acad. Sci. U. S. A.*, 1993, **90**, 5881–5888.
- 311 N. Ishida, K. Miyazaki, K. Kumagai and M. Rikimaru, *J. Antibiot.*, 1965, **18**, 68–76.
- 312 W. M. Maiese, M. P. Lechevalier, H. A. Lechevalier, J. Korshalla, N. Kuck, A. Fantini, M. J. Wildey, J. Thomas and M. Greenstein, *J. Antibiot.*, 1989, **42**, 558–563.
- 313 J. R. Baker, D. N. Woolfson, F. W. Muskett, R. G. Stoneman, M. D. Urbaniak and S. Caddick, *ChemBioChem*, 2007, **8**, 704–717.
- 314 K. V. Rao and W. P. Cullen, *Antibiot. Annu.*, 1959, **7**, 950–953.
- 315 M. Levine and M. Borthwick, *Virology*, 1963, **21**, 568–574.
- 316 A. Gupta and J. A. Imlay, *Proc. Natl. Acad. Sci. U. S. A.*, 2023, **120**, e2312110120.
- 317 C. B. Thorne and W. H. Peterson, *J. Biol. Chem.*, 1948, **176**, 413–428.
- 318 J. Bérddy, I. Horváth and A. Szentirmai, *Z. Allg. Mikrobiol.*, 1964, **4**, 232–235.
- 319 T. Nishimura, J. Kitajima, S. Omura and N. Tanaka, *J. Antibiot.*, 1981, **34**, 856–861.
- 320 H. Umezawa, S. Hayano, K. Maeda, Y. Ogata and Y. Okami, *Jpn. Med. J.*, 1950, **3**, 111–117.
- 321 N. Imamura, M. Nishijima, T. Takadera, K. Adachi, M. Sakai and H. Sano, *J. Antibiot.*, 1997, **50**, 8–12.
- 322 M. P. Singh, A. T. Menendez, P. J. Petersen, W. D. Ding, W. M. Maiese and M. Greenstein, *J. Antibiot.*, 1997, **50**, 785–787.
- 323 J. Kang, Y.-H. Cho and Y. Lee, *Microbiol. Spectr.*, 2022, **10**, e02312–e02322.
- 324 M. E. Flood, R. B. Herbert and F. G. Holliman, *J. Chem. Soc., Chem. Commun.*, 1970, 1514–1515.
- 325 M. Jancheva and T. Böttcher, *J. Am. Chem. Soc.*, 2021, **143**, 8344–8351.
- 326 M. B. Miller and B. L. Bassler, *Annu. Rev. Microbiol.*, 2001, **55**, 165–199.
- 327 R. Fernández-Piñar, M. Cámara, J.-F. Dubern, J. L. Ramos and M. Espinosa-Urgel, *Res. Microbiol.*, 2011, **162**, 773–781.
- 328 E. Déziel, F. Lépine, S. Milot, J. He, M. N. Mindrinos, R. G. Tompkins and L. G. Rahme, *Proc. Natl. Acad. Sci. U. S. A.*, 2004, **101**, 1339–1344.
- 329 F. S. Rossmann, T. Racek, D. Wobser, J. Puchalka, E. M. Rabener, M. Reiger, A. P. A. Hendrickx, A.-K. Diederich, K. Jung, C. Klein and J. Huebner, *PLoS Pathog.*, 2015, **11**, e1004653.
- 330 B. L. Bassler, M. Wright and M. R. Silverman, *Mol. Microbiol.*, 1994, **13**, 273–286.
- 331 X. Chen, S. Schauder, N. Potier, A. Van Dorsselaer, I. Pelczer, B. L. Bassler and F. M. Hughson, *Nature*, 2002, **415**, 545–549.
- 332 J. E. Silpe and B. L. Bassler, *Cell*, 2019, **176**, 268–280.
- 333 K. Papenfort, J. E. Silpe, K. R. Schramma, J.-P. Cong, M. R. Seyedsayamdost and B. L. Bassler, *Nat. Chem. Biol.*, 2017, **13**, 551–557.
- 334 Z. Erez, I. Steinberger-Levy, M. Shamir, S. Doron, A. Stokar-Avihail, Y. Peleg, S. Melamed, A. Leavitt, A. Savidor, S. Albeck, G. Amitai and R. Sorek, *Nature*, 2017, **541**, 488–493.
- 335 A. Dragoš, A. J. C. Andersen, C. N. Lozano-Andrade, P. J. Kempen, Á. T. Kovács and M. L. Strube, *Curr. Biol.*, 2021, **31**, 3479–3489.

