



ELSEVIER



Antibiotics: past, present and future

Matthew I Hutchings¹, Andrew W Truman² and Barrie Wilkinson²

The first antibiotic, salvarsan, was deployed in 1910. In just over 100 years antibiotics have drastically changed modern medicine and extended the average human lifespan by 23 years. The discovery of penicillin in 1928 started the golden age of natural product antibiotic discovery that peaked in the mid-1950s. Since then, a gradual decline in antibiotic discovery and development and the evolution of drug resistance in many human pathogens has led to the current antimicrobial resistance crisis. Here we give an overview of the history of antibiotic discovery, the major classes of antibiotics and where they come from. We argue that the future of antibiotic discovery looks bright as new technologies such as genome mining and editing are deployed to discover new natural products with diverse bioactivities. We also report on the current state of antibiotic development, with 45 drugs currently going through the clinical trials pipeline, including several new classes with novel modes of action that are in phase 3 clinical trials. Overall, there are promising signs for antibiotic discovery, but changes in financial models are required to translate scientific advances into clinically approved antibiotics.

Addresses

¹ School of Biological Sciences, University of East Anglia, Norwich Research Park, Norwich, NR4 7TJ, UK

² Department of Molecular Microbiology, John Innes Centre, Norwich Research Park, Norwich, NR4 7UH, UK

Corresponding authors: Hutchings, Matt (m.hutchings@uea.ac.uk), Truman, Andrew (andrew.truman@jic.ac.uk), Wilkinson, Barrie (barrie.wilkinson@jic.ac.uk)

Current Opinion in Microbiology 2019, 51:72–80

This review comes from a themed issue on **Antimicrobials**

Edited by **Matthew I Hutchings, Andrew W Truman and Barrie Wilkinson**

For a complete overview see the [Issue](#) and the [Editorial](#)

Available online 13th November 2019

<https://doi.org/10.1016/j.mib.2019.10.008>

1369-5274/© 2020 The Authors. Published by Elsevier Ltd. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

The development of antibiotics

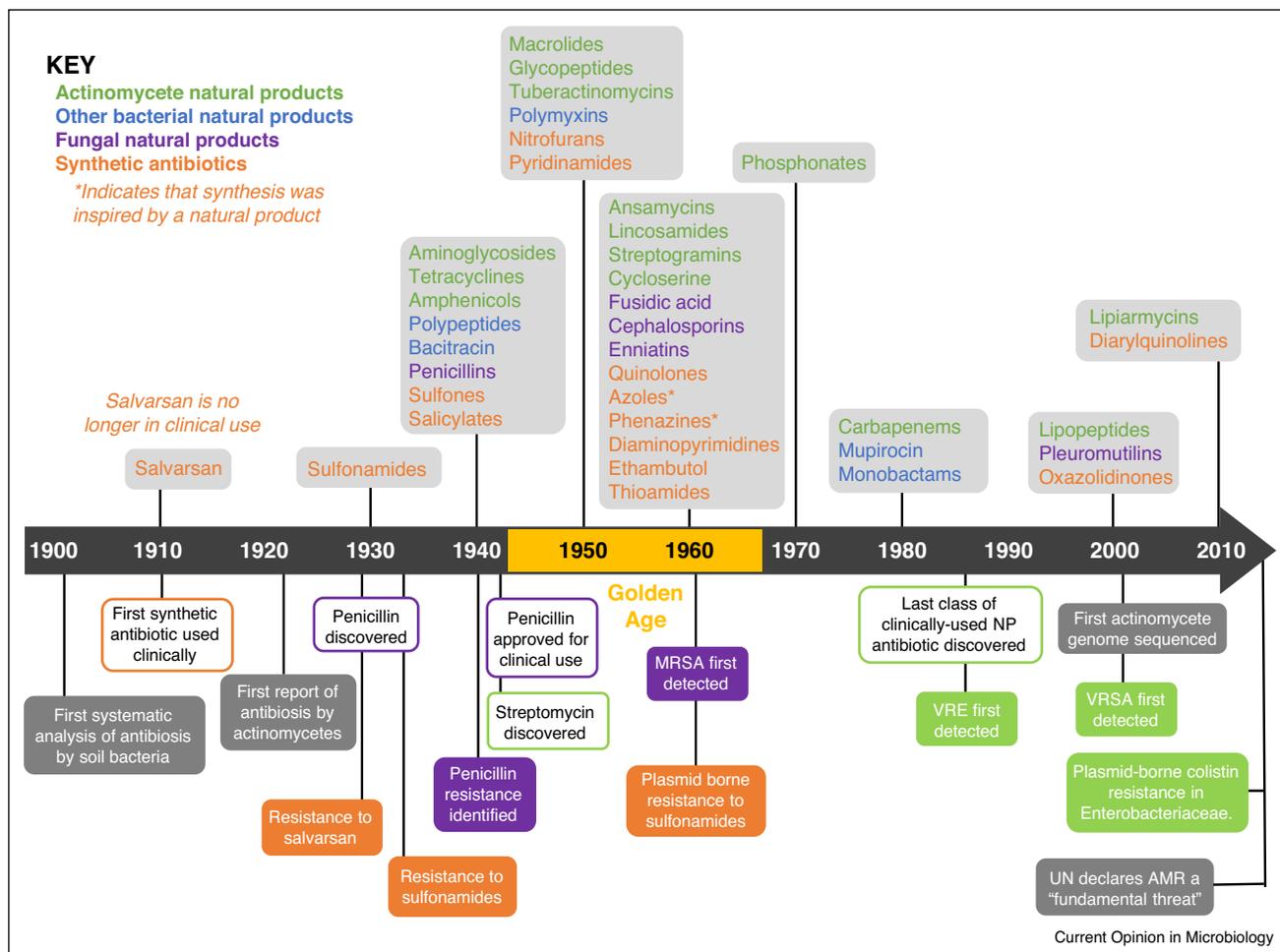
The introduction of antibiotics into clinical use was arguably the greatest medical breakthrough of the 20th century (Figure 1) [1]. In addition to treating infectious diseases, antibiotics made many modern medical procedures possible, including cancer treatment, organ transplants and open-heart surgery. However, misuse of these

valuable compounds has resulted in the rapid rise of antimicrobial resistance (AMR) with some infections now effectively untreatable [2]. The dangers of a post-antibiotic era has prompted policymakers to acknowledge this threat to human health and promise additional grant funding, which is gradually driving a resurgence of interest in antibiotic discovery and development [3]. The UK Government-commissioned O'Neill report predicted that without urgent action 10 million people a year will die from drug resistant infections by 2050 [4]. One of the key recommendations is to stimulate early stage drug discovery [4]. Given the relative lack of success in bringing effective synthetic antibiotics to the clinic [5], the best hope for developing a new generation of anti-infective drugs is to discover new microbial natural products (NPs) because these compounds are unrivalled in their chemical diversity and effectiveness as antibiotics [1]. Filamentous actinomycetes make 64% of the known NP antibiotic classes with the remainder made by other bacteria and fungi (Figure 2 and Table 1). Here we give a brief overview of the history of NP antibiotics and our prospects for discovering, developing and safeguarding a new generation of antibiotics.

A brief history of antibiotics

The use of antibiotic-producing microbes to prevent disease stretches back millennia, with traditional poultices of mouldy bread being used to treat open wounds in Serbia, China, Greece and Egypt more than 2000 years ago. The Eber's papyrus from 1550 BC is the oldest preserved medical document and includes mouldy bread and medicinal soil amongst its list of remedies [6]. An Anglo-Saxon recipe from 1000 years ago was also recently shown to kill MRSA (methicillin-resistant *Staphylococcus aureus*) [7*]. However, the development of anti-infective drugs and the underlying concept of chemotherapy is widely accredited to Paul Ehrlich, who developed the synthetic arsenic-based pro-drugs salvarsan (salvation arsenic) and neo-salvarsan circa 100 years ago to treat *Treponema pallidum*, the causative agent of syphilis [8] (Figure 1). This represented one of the first systematic screens for drug discovery using a library of synthetic compounds and was inspired by Ehrlich's work on dyes that specifically stained bacterial cells. Salvarsan was superseded by the sulfonamide prodrug Prontosil, discovered by Gerhard Domagk [9], a bacteriologist at Bayer who used the drug to save his daughter's arm from amputation. Domagk and colleagues were effectively continuing the work of Paul Ehrlich because the sulfa drugs were inspired by dyes that were used to selectively stain bacterial cells. Sulfonamides were the first truly effective, broad spectrum antimicrobials in clinical use

Figure 1



Timeline showing the decade new classes of antibiotic reached the clinic. The antibiotics are coloured per their source: green = actinomycetes, blue = other bacteria, purple = fungi and orange = synthetic. At the bottom of the timeline are key dates relating to antibiotic discovery and antimicrobial resistance, including the first reports of drug resistant strains methicillin-resistant *S. aureus* (MRSA), vancomycin-resistant enterococci (VRE), vancomycin-resistant *S. aureus* (VRSA) and plasmid-borne colistin resistance in Enterobacteriaceae.

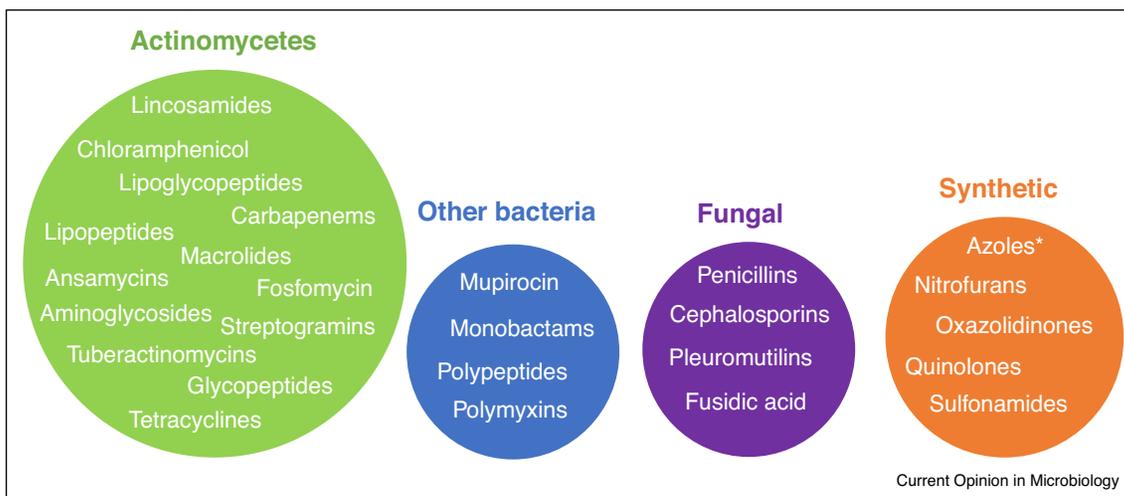
and are still in use today, but they were largely superseded by the discovery of penicillin, observed on a contaminated Petri dish by Alexander Fleming in 1928 [10]. Penicillin was later purified by Norman Heatley, Howard Florey, Ernst Chain and colleagues at Oxford, who were instrumental in the development of penicillin as a drug [11] (Figure 1). Dorothy Hodgkin solved the beta-lactam structure of penicillin in 1945 [12], resolving the famous debate between Robert Robinson, who favoured a thiazolidine-oxazolone structure, and several other notable chemists including Chain, Abrahams and Woodward, who believed it to be a beta-lactam [13]. This was an important breakthrough because it enabled the development of semi-synthetic derivatives to bypass penicillin resistance.

Antibiosis between microbes was described well before the discovery of penicillin, including by Louis Pasteur,

who proposed that microbes could secrete material to kill other bacteria [14]. The production of diffusible and heat-stable compounds by bacteria was being reported by the turn of the 20th century [15], and their utility in combatting infectious diseases had been explored. Arguably the first clinical use of an antibiotic was reported in the 1890s, where Emmerich and Löw used an extract of *Pseudomonas aeruginosa* (then known as *Bacillus pycyanus*) to treat hundreds of patients and this extract, called pyocyanase, was used until the 1910s [16]. Pyocyanase was active towards multiple pathogens and incorrectly believed to be an enzyme. Instead, the active components of pyocyanase was likely to be a mixture of pyocyanin, a quorum sensing phenazine, and 2-alkyl-4-hydroxy-quinolones [17].

The discoveries of penicillin, tyrocidine and numerous reports of the production of antimicrobial compounds by

Figure 2



Most clinically relevant classes of antibiotic are derived from natural products.

microorganisms, led Selman Waksman to start a systematic study of microbes as producers of antimicrobial compounds in the late 1930s. Waksman defined an antibiotic as ‘*a compound made by a microbe to destroy other microbes*’ and was instrumental in identifying soil-dwelling filamentous Actinomycetales (‘actinomycetes’) as prolific producers of antimicrobial compounds [18]. Waksman discovered numerous antibiotics made by soil-dwelling actinomycetes, including neomycin and streptomycin, the first agent active against tuberculosis [18]. Waksman’s pioneering work identified the genus *Streptomyces* as prolific producers of (natural products) NPs, or secondary metabolites, which are compounds not required for the normal growth, development, or reproduction of an organism in the laboratory. Many streptomycete NPs are active against bacteria, fungi, viruses, nematodes and insects and they have also been developed as anti-cancer and immunosuppressant drugs [19].

Waksman’s work initiated the Golden Age of antibiotic discovery from the 1940s to the 1960s. Most of these antibiotics are still in clinical use but their effectiveness has been eroded by the rise of AMR (Figure 1) [1]. In fact, the rapid and relatively easy discovery of multiple classes (and variations therein) of NP antibiotics during a relatively short period led to the excessive use of these drugs. This, coupled with a faltering antibiotic discovery pipeline from the 1970s onwards, has led to the current situation with few new antibiotics in the clinical trials pipeline [1]. Hence, most antibiotics in clinical trials today are derivatives of known classes of NP or synthetic antibiotics rather than new classes of antibiotic (Table S1). Notably, this hiatus in antibiotic discovery aligns with a decline in the discovery of new NP families and the persistent rediscovery of known compounds in

screening campaigns using microbial, and predominantly actinomycete, fermentation extracts [1]. This, in part, led to a belief that all the ‘low-hanging fruit’ had been harvested and resulted in most of major pharmaceutical and agrochemical companies shutting down their NP discovery departments.

The divestment in NP research was accompanied by an investment in numerous high-throughput screening (HTS) programmes that aimed to discover new synthetic antibiotics, but these have proved unsuccessful. For example, 70 HTS campaigns were conducted by GlaxoSmithKline (GSK) over seven years using a collection of approximately 500 000 compounds, but this yielded very few leads, and no candidates for development [20]. Similarly, 65 HTS campaigns by AstraZeneca provided a few leads but none that were active against multi-drug resistant Gram-negative bacteria [21]. In recent years however, the discovery of new antibiotic-producing strains in under-explored environments combined with new tools for genome mining has reinvigorated the NP discovery field, for example [22,23,24].

Why do microorganisms make antibiotics?

Of all the antibiotics discovered between 1945 and 1978, 55% came from the genus *Streptomyces* (Figure 1) [25]. Several theories have been proposed to explain why soil microbes make so many bioactive NPs. The most likely explanation is that they have multiple functions, acting as chemical weapons to kill competitors in the soil either as protection (defensive) or predation (offensive), as signalling molecules to close relatives or to mediate interactions with eukaryotic hosts such as insects and plants [26–28]. This is consistent with evidence that *Streptomyces* species and other filamentous actinomycetes evolved circa

Table 1

All classes of clinically used antibiotics and their source

Class ^a	Discovery reported ^b	Introduced clinically	Example (and producing organism)	Molecular target
Antibiotics from actinomycetes				
Aminoglycosides	1944	1946	Kanamycin A (<i>Streptomyces kanamyceticus</i>)	Protein synthesis: 30S ribosomal subunit
Tetracyclines	1948	1948	Tetracycline (<i>Streptomyces aureofaciens</i>)	Protein synthesis: 30S ribosomal subunit
Amphenicols	1947	1949	Chloramphenicol (<i>Streptomyces venezuelae</i>)	Protein synthesis: 50S ribosomal subunit
Macrolides	1952	1952	Erythromycin (<i>Saccharopolyspora erythraea</i>)	Protein synthesis: 50S ribosomal subunit
Tuberactinomycins	1951	1953	Viomycin (<i>Streptomyces puniceus</i>)	Protein synthesis: 30S and 50S ribosomal subunits (binds to the intersubunit bridge B2a)
Glycopeptides	1954	1958	Vancomycin (<i>Amycolatopsis orientalis</i>)	Cell wall synthesis: D-Ala-D-Ala termini of lipid II
Lincosamides	1962	1963	Clindamycin Semi-synthetic derivative of lincomycin (<i>Streptomyces lincolnensis</i>)	Protein synthesis: 50S ribosomal subunit
Ansamycins	1959	1963	Rifamycin SV Semi-synthetic derivative of rifamycin (<i>Amycolatopsis rifamycinica</i>)	Nucleic acid synthesis: RNA polymerase
Cycloserines	1955	1964	Seromycin (<i>Streptomyces orchidaceus</i>)	Cell wall synthesis: inhibition of alanine racemase and D-alanine-D-alanine ligase
Streptogramins	1953	1965	Pristinamycin (<i>Streptomyces pristinaespiralis</i>)	Protein synthesis: 50S ribosomal subunit
Phosphonates	1969	1971	Fosfomycin (<i>Streptomyces fradiae</i>)	Cell wall synthesis: MurA (UDP-GlcNAc-3-enolpyruvyltransferase) inhibition
Carbapenems	1976	1985	Meropenem Synthetic molecule based on thienamycin (<i>Streptomyces cattleya</i>)	Cell wall synthesis: penicillin-binding proteins
Lipopeptides	1987	2003	Daptomycin (<i>Streptomyces roseosporus</i>)	Cell wall: cell membrane disruption.
Lipiamycins	1975	2011	Fidaxomicin (<i>Dactylosporangium aurantiacum</i> subsp. <i>hamdenensis</i>)	Nucleic acid synthesis: RNA polymerase
Antibiotics from other bacteria				
Polypeptides	1939	1941	Gramicidin A (<i>Bacillus brevis</i>)	Cell wall: forms ion channels that increase the permeability of the bacterial cell membrane
Bacitracin	1945	1948	Bacitracin A (<i>Bacillus subtilis</i>)	Cell wall synthesis: inhibition of dephosphorylation of C ₅₅ -isoprenyl pyrophosphate
Polymyxins	1950	1959	Colistin (<i>Paenibacillus polymyxa</i>)	Cell wall: cell membrane disruption
Mupirocin	1971	1985	Mupirocin (<i>Pseudomonas fluorescens</i>)	Protein synthesis: isoleucyl t-RNA synthetase
Monobactams	1981	1986	Aztreonam Synthetic molecule based on SQ 26,180 (<i>Chromobacterium violaceum</i>)	Cell wall synthesis: penicillin-binding proteins
Antibiotics from fungi				
Penicillins	1929	1943	Amoxicillin Semi-synthetic derivative of penicillin (<i>Penicillium chrysogenum</i>)	Cell wall synthesis: penicillin-binding proteins
Fusidic acid	1958	1962	Fusidic acid (<i>Fusidium coccineum</i>)	Protein synthesis: elongation factor G
Enniatins ^c	1953	1963	Fusafungine (<i>Fusarium lateritium</i>)	Cell wall: cell membrane disruption
Cephalosporins	1948	1964	Cefacetrile Semi-synthetic derivative of cephalosporin C (<i>Acremonium chrysogenum</i>)	Cell wall synthesis: penicillin-binding proteins
Pleuromutilins	1951	2007	Retapamulin Semi-synthetic derivative of pleuromutilin (<i>Pleurotus mutilus</i>)	Protein synthesis: 50S ribosomal subunit
Synthetic antibiotics				
Arsphenamines ^d	1907	1910	Salvarsan	Not known

Table 1 (Continued)

Class ^a	Discovery reported ^b	Introduced clinically	Example (and producing organism)	Molecular target
Sulfonamides	1932	1936	Mafenide	Folate synthesis: inhibition of dihydropteroate synthetase
Salicylates ^e	1902	1943	4-Aminosalicylic acid	Folate synthesis: prodrug that inhibits dihydrofolate reductase
Sulfones	1908	1945	Dapsone	Folate synthesis: inhibition of dihydropteroate synthetase
Pyridinamides	1952	1952	Isoniazid	Cell wall: prodrug that inhibits the synthesis of mycolic acids
Nitrofurans	1945	1953	Nitrofurantoin	DNA synthesis: DNA damage
Azoles ^f	1959	1960	Metronidazole	DNA synthesis: DNA damage
(Fluoro)quinolones	1962	1962	Ciprofloxacin	DNA synthesis: inhibition of DNA gyrase, and topoisomerase IV
Diaminopyrimidines	1950	1962	Trimethoprim	Folate synthesis: inhibition of dihydrofolate reductase
Ethambutol	1962	1962	Ethambutol	Cell wall: arabinosyl transferase inhibition
Thioamides	1956	1965	Ethionamide	Cell wall: prodrug that inhibits the synthesis of mycolic acids
Phenazines ^f	1954	1969	Clofazimine	DNA synthesis: binds to guanine bases
Oxazolidinones	1987	2000	Linezolid	Protein synthesis: 50S ribosomal subunit
Diarylquinolines	2004	2012	Bedaquiline	ATP synthesis: proton pump inhibition

^a Classes are defined by origin, structure and/or mechanism of action, which distinguishes between bacitracin, colistin and daptomycin, for example.

^b Year reported refers to first report in literature.

^c The European Medicines Agency recommended the withdrawal of fusafungine from the market in February 2016.

^d Salvarsan is no longer in clinical use.

^e Salicylic acids are found in nature, but this was not the source of this class of antibiotic.

^f Compound synthesis was inspired by natural antibiotic classes.

440 million years ago, around the same time that plants first colonized land [25,29]. The filamentous growth of these bacteria would have provided an advantage in colonizing plant roots and we speculate that many of their NPs may have evolved or been co-opted to mediate these interactions [30].

One of the more surprising discoveries to arise from microbial genome sequencing is that many bacteria and fungi encode many more NP pathways than they actually make in the laboratory [31*]. In general, at least three quarters of their potential NP capability is not switched on *in vitro* and this discovery has triggered huge efforts to develop tools and techniques to activate their “cryptic” biosynthetic gene clusters (BGCs) in the hope of discovering novel chemical scaffolds with useful bioactivities [32,33**,34,35,36**]. Many studies have demonstrated that when activated or expressed heterologously, silent BGCs encode functional NP biosynthetic pathways [34]. This suggests that production of these compounds is triggered by environmental cues or by host organisms. Many invertebrates, including insects and marine sponges, form defensive and mutually beneficial symbioses (defensive mutualisms) with antibiotic-producing bacteria and it seems likely that most if not all land plants do the same [26,37–39]. Studying these bacteria in the context of their host using advanced techniques, such as stable isotope probing (SIP) and imaging mass

spectrometry (IMS), may be one way to identify the thousands of novel compounds encoded by silent BGCs and to identify the NPs that are most important to their hosts [40].

Prospects for natural product antibiotic discovery

In the Golden Age of antibiotic discovery, new antibiotic classes were being discovered on an almost yearly basis by isolation of likely antibiotic-producing organisms from soil samples. However, a finite number of NP classes from easy-to-cultivate bacteria meant that compound rediscovery soon became a problem (Figure 1). More recently, the NP discovery field has been reinvigorated by the discovery of new antibiotic-producing strains in under-explored environments, combined with new tools for genome mining and heterologous pathway expression.

Under-explored environments and ecological niches

It is now clear that only a tiny fraction of the soils on earth have been sampled for antibiotic producers. Sampling more widely is likely to yield numerous new strains and BGCs, even from this traditional sampling environment. In addition, sampling under-explored environments that were inaccessible or unknown during the Golden Age is yielding new chemical structures [20,37,38]. These include the marine environment, where the marine actinomycete genus *Salinospora* has proven to be a source of

multiple structurally novel NPs [41] such as salinosporamide A (Marizomib), which has anticancer activity and is currently in Phase III clinical trials for the treatment of glioblastoma [42].

Mutualistic co-evolved bacteria might also be an excellent source of new NPs and studying these niches has the added advantage of uncovering interesting underlying biology and the opportunity to understand what these molecules actually do in nature [43**,44**]. Bacterial symbionts of marine invertebrates such as sponges are a rich source of novel NPs. For example, *Candidatus Entotheonella* species are uncultivated symbionts of the marine sponge *Theonella swinhoei* [45] and were shown to produce almost all the bioactive polyketides and modified peptides isolated from a chemotype of *T. swinhoei*.

Sequencing of the human microbiome has also revealed many NP BGCs across Actinobacteria and other bacterial phyla, and the antibiotic lactocillin was identified from a human vaginal isolate [46]. Another antibacterial compound, lugdunin, was isolated from the commensal nasal bacterium *Staphylococcus lugdunensis* which prohibits colonization by *S. aureus* and is active in animal models, with a high barrier to the development of resistance [47].

Difficult to cultivate bacteria

Genomic data suggesting the presence of novel BGCs in *Clostridium* bacteria prompted Hertweck and colleagues to investigate the antibiotic-producing potential of this genus, as no NPs had been characterised from clostridia. *Clostridium cellulolyticum* grown under standard laboratory conditions yielded no NPs, so fermentation was repeated with added aqueous soil extracts, as the bacterium had been isolated from decayed grass compost. This triggered the production of closthioamide, a new class of polythioamide antibiotic [48]. In another elegant example, the antibiotic humimycin was discovered by synthesising a putative peptide NP that was bioinformatically predicted from the genome of the actinomycete *Rhodococcus equi*, an opportunist human pathogen [23**].

Other novel approaches have included the isolation of hard to culture bacteria from soil using diffusion chambers that allow for the growth of the pure bacterium in a complex natural environment [49]. This was miniaturised into an isolation chip (iChip) and used to culture 10 000 soil isolates that were otherwise intractable to laboratory fermentation. Extracts generated from these were then screened for antimicrobial activity and one resulted in the identification of the antibacterial peptide teixobactin that is produced by *Eleftheria terrae* [50]. These discoveries and recent metagenomics studies [51] highlight the continued relevance of traditional soil environments for antibiotic discovery.

The development of improved sampling methodologies for under-explored environments and difficult to cultivate bacteria, combined with new genetic tools and technologies to activate interesting BGCs, is likely to lead to the discovery of thousands of new bioactive compounds over the next 20 years. It is highly probable that some fraction of these will form the basis of new anti-infectives for clinical medicine, although this will require improved financial models to incentivise the development of new antibiotics.

Prospects for clinical development

As of December 2018, there are 45 new antibiotic candidates in clinical trials for the US market (Table S1) [52]. Of these, 28 belong to known NP classes while 17 are synthetic and comprise 12 classes, of which seven are new. The NP classes include 13 based on beta-lactams, which was the first class of NP antibiotic to be discovered back in 1928 (Figure 1). Five of these are variant beta-lactams, two are hybrids (to a glycopeptide and a siderophore) and seven are combinations with beta-lactamase inhibitors (Table S1). There are five new tetracyclines, a class which was first described in 1945 and introduced into the clinic in 1948, an aminoglycoside (1943), a distamycin (1962), a fusidane (1945), a macrolide (1952), a pleuromutilin (1950) and two polymyxins (1947). The fusidane (fusidic acid) is a fungal NP which is in Phase III trials in the United States, but it has already been used clinically elsewhere in the world. There are two new synthetic classes in Phase III clinical trials: ridinilazole, which specifically blocks cell division in *Clostridium difficile* through a mechanism that has not been revealed; and murepavedin, which has a novel mechanism of action, inhibiting LptD to block lipopolysaccharide transport to the outer membrane [53]. Murepavidin is effective against drug resistant *P. aeruginosa*, one of the hardest pathogens to treat, particularly in patients with cystic fibrosis. It is also encouraging that four of the nine compounds in Phase II clinical trials represent novel classes, but this is still a modest number for the therapeutic area and is insufficient to combat multidrug-resistant Gram-negative pathogens given the historically high attrition rate for compounds making it through clinical trials to clinical utility. AntibioticDB is an open access database that records candidate antibiotics, including antibiotics under pre-clinical development, those in clinical trials and discontinued drugs [54].

Unfortunately, most of the large pharmaceutical companies have left the field of NP discovery, and this work is now chiefly undertaken by academic labs and small to medium-sized companies. Only two of the 45 drugs currently in development belong to big pharmaceutical companies: the synthetic gepotacidin inhibits topoisomerase II through a mechanism distinct from that of quinolones and is being developed by GSK to treat gonorrhoea (phase 2) while Merck have a beta-lactam/lactamase combination in phase III clinical trials. The most notable NP antibiotic success in recent years was the introduction of Cubicin (daptomycin) onto the market by

Cubist in 2003, and sales of this drug are now more than \$1 billion a year. Cubicin is used by injection to treat vancomycin-resistant *S. aureus* (VRSA) and was discovered from *Streptomyces roseosporus* in 1987. In 2011, Cubist also purchased Optimer Pharmaceuticals, who secured clinical approval for Dificid (fidaxomicin, produced by the rare actinomycete *Dactyloporangium auranticus* subsp. *hamdenensis*). This is the newest NP class to be introduced into the clinic, despite being discovered in 1975, before daptomycin [55]. Merck purchased Cubist in 2015 for \$9.5B but have since closed the discovery arm of Cubist, which was heavily involved in NP discovery.

In 2014 Sanofi and Fraunhofer announced the creation of a NP Centre of Excellence with the goal of identifying novel compounds to accelerate the discovery and development of new antibiotics. In 2016 Sanofi further announced a partnership with Warp Drive Bio to collaborate on the development of novel oncology therapies and antibiotics by using next generation sequencing and genome mining (on a massive scale) to identify new NPs but this ended in 2017. In 2018 Warp Drive Bio was effectively merged with Revolution Medicines, which is now focussed on oncology rather than anti-infectives, although Warp Drive Bio's genome mining platform has recently been acquired by Ginkgo Bioworks. Roche have several strategic alliances, such as with Spero, which currently has two antibiotics in phase I and another in phase III clinical trials (Table S1). Several companies, including Genentech, are working on antibody-antibiotic conjugates (AACs) [56]. Of the larger to mid-sized companies, Basilea is a major active player and focuses on the development of innovative antibiotics, antifungals and oncology drugs. In addition, there are innovative small to midsized companies in the antibacterial and antifungal discovery space including Tetrphase Pharma, which currently has two antibiotics in phase I clinical trials and recently had two more approved for use (Table S1). There is a heavy NP influence on all these companies, which appear to be using semisynthetic or total synthesis approaches within very specific areas of chemical space around known NPs such as polyenes, macrolides and tetracyclines.

Beyond the scientific difficulties associated with antibiotic discovery and development, there are a plethora of regulatory, economic, business and societal issues that must be addressed in order to protect and maximise the potential of our existing and future arsenal of clinical agents, while at the same time promoting the investment and culture changes required to invigorate antibiotics R&D to meet the challenges raised by AMR [57,58]. These have been analysed and recommendations made in several key reports including those by O'Neill and the Pew Trust [4,52]. Mossialos and colleagues comprehensively reviewed 47 incentive strategies for the development of new antibiotics and concluded that a framework of multiple incentives and policies is required [59].

Summary and outlook

The rise in bacterial infections that are resistant to almost all known antibiotics is alarming, yet it is only in the last few years that governments have begun to tackle this problem seriously. This global wake-up call has stimulated a debate about how best to combat AMR and prompted the UK government to appoint an economist, Lord Jim O'Neill, to lead a strategic review [4]. The appointment of an economist highlighted the complexities of bringing to market a drug that, if functionally successful, will be dosed for only a short time. Combined with historically low prices, and the likelihood that any new antibiotic with a unique mode of action will most probably be restricted as a treatment of last resort, the economics of antibiotic R&D is a major disincentive to investment. To address these problems innovative solutions are required that provide a reimbursement model that delinks revenue from drug sales.

Scientifically, the identification of new chemical matter with the unique physicochemical characteristics required for antibiotic discovery and development is a key challenge. NPs still represent the most likely source of new materials given the advances described in this review. Even the best-studied antibiotic producers, the streptomycetes, have been vastly under-sampled in terms of their capability, and there is confidence from the study of organisms from underexploited environments, ecological considerations, and genome sequencing that thousands of NP antibiotics await discovery across the bacterial kingdom. New tools and techniques such as CRISPR/Cas9-mediated genome editing are available to exploit these observations, although there is no universal strategy for the expression of silent BGCs. Recent advances have led to the discovery of many new molecular structures with exceptional biological activities [34], and further advances in this area will undoubtedly accelerate this rate of discovery.

Thus, governments are starting to act and there is much to be optimistic about, not least the fact that most of the NP antibiotics that have been discovered come from a small fraction of the microbes on Earth. With suitable global action, this should lead to a renewed antibiotic pipeline to combat AMR alongside other emergent technologies, such as vaccines, antibody-antibiotic conjugates, probiotics, phage therapy and rapid diagnostics [60].

Competing interest statement

Nothing declared.

Disclosure Statement

Given their role as Guest Editor, Matt Hutchings, Andrew Truman, Barrie Wilkinson, had no involvement in the peer-review of this article and has no access to information regarding its peer-review. Full responsibility

for the editorial process for this article was delegated to one of the Editors in Chief.

Acknowledgements

We thank our colleagues on the Norwich Research Park for stimulating discussions. We acknowledge financial support for our work on NP antibiotics through PhD studentships from the Biotechnology and Biological Sciences Research Council (BBSRC Doctoral Training Program grant BB/M011216/1) and the Natural Environment Research Council (NERC Doctoral Training Programme grant NE/L002582/1), responsive mode grants from BBSRC, MRC and NERC, and a Royal Society University Research Fellowship for AWT.

Appendix A. Supplementary data

Supplementary material related to this article can be found, in the online version, at doi:<https://doi.org/10.1016/j.mib.2019.10.008>.

References and recommended reading

Papers of particular interest, published within the period of review, have been highlighted as:

- of special interest
- of outstanding interest

1. Katz L, Baltz RH: **Natural product discovery: past, present, and future.** *J Ind Microbiol Biotechnol* 2016, **43**:155-176.
 2. Prescott JF: **The resistance tsunami, antimicrobial stewardship, and the golden age of microbiology.** *Vet Microbiol* 2014, **171**:273-278.
 3. Walsh CT, Wenczewicz TA: **Prospects for new antibiotics: a molecule-centered perspective.** *J Antibiot* 2014, **67**:7-22.
 4. Neil JO: *Report on Antimicrobial Resistance.* 2016 <https://amr-review.org>.
 5. Payne DJ, Gwynn MN, Holmes DJ, Pompliano DL: **Drugs for bad bugs: confronting the challenges of antibacterial discovery.** *Nat Rev Drug Discovery* 2006, **6**:29-40.
 6. Haas LF: **Papyrus of Ebers and Smith.** *J Neurol Neurosurg Psychiatr* 1999, **67**:572-578.
 7. Harrison F, Roberts AEL, Gabriliska R, Rumbaugh KP, Lee C, Diggle SP: **A 1,000-year-old antimicrobial remedy with antistaphylococcal activity.** *mBio* 2015, **6**:e01129
- This paper follows an AngloSaxon recipe to make an antimicrobial substance that can inhibit MRSA.
8. Gelpi A, Gilbertson A, Tucker JD: **Magic bullet: Paul Ehrlich, Salvarsan and the birth of venereology.** *Sex Transm Infect* 2015, **91**:68-69.
 9. Otten H: **Domagk and the development of the sulphonamides.** *J Antimicrobial Chemother* 1986, **17**:689-690.
 10. Fleming A: **On the antibacterial action of cultures of a penicillium, with special reference to their use in the isolation of *B. influenzae*.** *Br J Exp Pathol* 1929, **10**:226-236.
 11. Abraham EP, Chain E, Fletcher CM, Gardner AD, Heatley NG, Jennings MA, Florey HW: **Further observations on penicillin.** *Lancet* 1941, **238**:177-189.
 12. Hodgkin DC: **X-ray crystallographic investigation of the structure of penicillin.** *Adv Sci* 1949, **6**:85-89.
 13. Curtis R, Jones J: **Robert Robinson and penicillin: an unnoticed document in the saga of its structure.** *J Pept Sci* 2007, **13**:769-775.
 14. Brunel J: **Antibiosis from Pasteur to Fleming.** *J Hist Med Allied Sci* 1951, **6**:287-301.
 15. Frost WD: **The antagonism exhibited by certain saprophytic bacteria against the *Bacillus typhosus*.** *J Infect Dis* 2016, **1**:1-44.
 16. Emmerich R, Löw O: **Bakteriolytische Enzyme als Ursache der erworbenen Immunität und die Heilung von Infektionskrankheiten durch dieselben.** *Zeitschrift für Hygiene und Infektionskrankheiten* 1899, **31**:1-65.
 17. Hays EE, Wells IC, Katzman PA, Cain CK, Jacobs FA, Thayer SA, Doisy EA, Gaby WL, Roberts EC, Muir RD *et al.*: **Antibiotic substances produced by *Pseudomonas aeruginosa*.** *J Biol Chem* 1945, **159**:725.
 18. Waksman SA, Schatz A, Reynolds DM: **Production of antibiotic substances by actinomycetes.** *Ann N Y Acad Sci* 2010, **1213**:112-124.
 19. Newman DJ, Cragg GM: **Natural products as sources of new drugs from 1981 to 2014.** *J Nat Prod* 2016, **79**:629-661.
 20. Payne DJ, Miller LF, Findlay D, Anderson J, Marks L: **Time for a change: addressing R&D and commercialization challenges for antibacterials.** *Philos Trans R Soc Lond B Biol Sci* 2015, **370**.
 21. Tommasi R, Brown DG, Walkup GK, Manchester JI, Miller AA: **ESKAPEing the labyrinth of antibacterial discovery.** *Nat Rev Drug Discov* 2015, **14**:529-542.
 22. Qin Z, Munnoch JT, Devine R, Holmes NA, Seipke RF, Wilkinson KA, Wilkinson B, Hutchings MI: **Formicamycins, antibacterial polyketides produced by *Streptomyces formicae* isolated from African Tetraponera plant-ants.** *Chem Sci* 2017, **8**:3218-3227.
 23. Chu J, Vila-Farres X, Inoyama D, Ternei M, Cohen LJ, Gordon EA, Reddy BVB, Charlop-Powers Z, Zebroski HA, Gallardo-Macias R *et al.*: **Discovery of MRSA active antibiotics using primary sequence from the human microbiome.** *Nat Chem Biol* 2016, **12**:1004-1006
- This paper reports on a new technique to make synthetic natural products after predicting their structures through sequence analysis. They used it to discover the humimycins.
24. Luo Y, Huang H, Liang J, Wang M, Lu L, Shao Z, Cobb RE, Zhao H: **Activation and characterization of a cryptic polycyclic tetramate macrolactam biosynthetic gene cluster.** *Nat Commun* 2013, **4**:2894.
 25. Embley TM: **The molecular phylogeny and systematics of the actinomycetes.** *Annu Rev Microbiol* 1994, **48**:257-289.
 26. Seipke RF, Kaltenpoth M, Hutchings MI: ***Streptomyces* as symbionts: an emerging and widespread theme?** *FEMS Microbiol Rev* 2012, **36**:862-876.
 27. Klassen JL: **Microbial secondary metabolites and their impacts on insect symbioses.** *Curr Opin Insect Sci* 2014, **4**:15-22.
 28. Traxler MF, Kolter R: **Natural products in soil microbe interactions and evolution.** *Nat Prod Rep* 2015, **32**:956-970.
 29. Wellman CH, Osterloff PL, Mohiuddin U: **Fragments of the earliest land plants.** *Nature* 2003, **425**:282-285.
 30. van der Meij A, Worsley SF, Hutchings MI, van Wezel GP: **Chemical ecology of antibiotic production by actinomycetes.** *FEMS Microbiol Rev* 2017, **41**:392-416.
 31. Blin K, Shaw S, Steinke K, Villebro R, Ziemert N, Lee SY, Medema M, Weber T: **antiSMASH 5.0: updates to the secondary metabolite genome mining pipeline.** *Nucl Acids Res* 2019, **47**:W81-87
- An update on the invaluable antiSMASH software which can predict secondary metabolite gene clusters from genome sequences.
32. Bauman KD, Li J, Murata K, Mantovani SM, Daheh S, Nizet V, Luhavaya H, Moore BS: **Refactoring the cryptic streptophenazine biosynthetic gene cluster unites phenazine, polyketide, and nonribosomal peptide biochemistry.** *Cell Chem Biol* 2019, **26**:724-736.e7.
 33. Moon K, Xu F, Zhang C, Seyedsayamdost MR: **Bioactivity-HiTES unveils cryptic antibiotics encoded in actinomycete bacteria.** *ACS Chem Biol* 2019, **14**:767-774
- This paper reports on a new high throughput method for screening elicitors of cryptic secondary metabolites and they use it to discover a novel molecule.

34. Rutledge PJ, Challis GL: **Discovery of microbial natural products by activation of silent biosynthetic gene clusters.** *Nat Rev Microbiol* 2015, **13**:509-523.
35. Zhang X, Hindra, Elliot MA: **Unlocking the trove of metabolic treasures: activating silent biosynthetic gene clusters in bacteria and fungi.** *Curr Opin Microbiol* 2019, **51**:9-15.
36. Gehrke EJ, Zhang X, Pimentel-Elardo SM, Johnson AR, Rees CA, Jones SE, Hindra, Gehrke SS, Turvey S, Boursalie S *et al.*: **Silencing cryptic specialized metabolism in *Streptomyces* by the nucleoid-associated protein Lsr2.** *eLife* 2019, **8**:W202
- This paper reports that by deleting the nucleoid-associated protein gene *lsr2* it is possible to switch on cryptic BGCs in diverse *Streptomyces* strains because *Lsr2* binds preferentially to secondary metabolite BGCs.
37. Kaltenpoth M: **Actinobacteria as mutualists: general healthcare for insects?** *Trends Microbiol* 2009, **17**:529-535.
38. Palaniyandi SA, Yang SH, Zhang L, Suh J-W: **Effects of actinobacteria on plant disease suppression and growth promotion.** *Appl Microbiol Biotechnol* 2013, **97**:9621-9636.
39. Viaene T, Langendries S, Beirinckx S, Maes M, Goormachtig S: ***Streptomyces* as a plant's best friend?** *FEMS Microbiol Ecol* 2016, **92**:fiw119-11.
40. Hoefler BC, Stubbendieck RM, Josyula NK, Moisan SM, Schulze EM, Straight PD: **A link between linear mycin biosynthesis and extracellular vesicle genesis connects specialized metabolism and bacterial membrane physiology.** *Cell Chem Biol* 2017, **24**:1238-1249.e7.
41. Ziemert N, Lechner A, Wietz M, Millán-Aguiñaga N, Chavarria KL, Jensen PR: **Diversity and evolution of secondary metabolism in the marine actinomycete genus *Salinispora*.** *Proc Natl Acad Sci U S A* 2014, **111**:E1130-E1139.
42. Gulder TAM, Moore BS: **Salinosporamide natural products: potent 20 S proteasome inhibitors as promising cancer chemotherapeutics.** *Angew Chem Int Ed Engl* 2010, **49**:9346-9367.
43. Heine D, Holmes NA, Worsley SF, Santos ACA, Innocent TM, Jensen PR, Scherlach K, Patrick EH, Yu DW, Murrell JC, Vieria PC *et al.*: **Chemical warfare between leafcutter ant symbionts and a co-evolved pathogen.** *Nat Commun* 2018, **9**:2208
- This paper reports that a specialised pathogen of leafcutter ants makes secondary metabolites that inhibit the growth of the ant defensive mutualist bacteria and also modify the grooming behaviour of the ants.
44. Zan J, Li Z, Tianero MD, Davis J, Hill RT, Donia MS: **A microbial factory for defensive kahalalides in a tripartite marine symbiosis.** *Science* 2019, **364**:eaaw6732
- This paper reports on a novel bacterial symbiont of a marine alga which makes a group of potent antibiotics as defence molecules for its host using amino acids provided by the host.
45. Wilson MC, Mori T, Rückert C, Uriá AR, Helf MJ, Takada K, Gernert C, Steffens UAE, Heycke N, Schmitt S *et al.*: **An environmental bacterial taxon with a large and distinct metabolic repertoire.** *Nature* 2015, **506**:58-62.
46. Donia MS, Cimermancic P, Schulze CJ, Wieland Brown LC, Martin J, Mitreva M, Clardy J, Linington RG, Fischbach MA: **A systematic analysis of biosynthetic gene clusters in the human microbiome reveals a common family of antibiotics.** *Cell* 2014, **158**:1402-1414.
47. Zipperer A, Konnerth MC, Laux C, Berscheid A, Janek D, Weidenmaier C, Burian M, Schilling NA, Slavetinsky C, Marschal M *et al.*: **Human commensals producing a novel antibiotic impair pathogen colonization.** *Nature* 2016, **535**:511-516.
48. Lincke T, Behnken S, Ishida K, Roth M, Hertweck C: **Closthioamide: an unprecedented polythioamide antibiotic from the strictly anaerobic bacterium *Clostridium cellulolyticum*.** *Angew Chem Int Ed Engl* 2010, **49**:2011-2013.
49. Nichols D, Cahoon N, Trakhtenberg EM, Pham L, Mehta A, Belanger A, Kanigan T, Lewis K, Epstein SS: **Use of ichip for high-throughput in situ cultivation of "uncultivable" microbial species.** *Appl Environ Microbiol* 2010, **76**:2445-2450.
50. Ling LL, Schneider T, Peoples AJ, Spoering AL, Engels I, Conlon BP, Mueller A, Schäberle TF, Hughes DE, Epstein S *et al.*: **A new antibiotic kills pathogens without detectable resistance.** *Nature* 2015, **517**:455-459.
51. Charlop-Powers Z, Pregitzer CC, Lemetre C, Ternei MA, Maniko J, Hover BM, Calle PY, McGuire KL, Garbarino J, Forgione HM *et al.*: **Urban park soil microbiomes are a rich reservoir of natural product biosynthetic diversity.** *Proc Natl Acad Sci U S A* 2016, **113**:14811-14816.
52. PEW Trust: **Antibiotics Currently in Global Clinical Development.** 2019 <https://www.pewtrusts.org>.
53. Srinivas N, Jetter P, Ueberbacher BJ, Werneburg M, Zerbe K, Steinmann J, Van der Meijden B, Bernardini F, Lederer A, Dias RLA *et al.*: **Peptidomimetic antibiotics target outer-membrane biogenesis in *Pseudomonas aeruginosa*.** *Science* 2010, **327**:1010-1013.
54. Farrell LJ, Lo R, Wanford JJ, Jenkins A, Maxwell A, Piddock LVJ: **Revitalizing the drug pipeline: AntibioticDB, an open access database to aid antibacterial research and development.** *J Antimicrob Chemother* 2018, **73**:2284-2297.
55. Jarrad AM, Karoli T, Blaskovich MAT, Lyras D, Cooper MA: ***Clostridium difficile* drug pipeline: challenges in discovery and development of new agents.** *J Med Chem* 2015, **58**:5164-5185.
56. Lehar SM, Pillow T, Xu M, Staben L, Kajihara KK, Vandlen R, DePalatis L, Raab H, Hazenbos WL, Morisaki JH *et al.*: **Novel antibody-antibiotic conjugate eliminates intracellular *S. aureus*.** *Nature* 2015, **527**:323-328.
57. van der Meer JWM, Fears R, Davies SC, Meulen ter V: **Antimicrobial innovation: combining commitment, creativity and coherence.** *Nat Rev Drug Discov* 2014, **13**:709-710.
58. Mullard A: **Momentum builds around new antibiotic business models.** *Nat Rev Drug Discov* 2014, **13**:711-713.
59. Renwick MJ, Brogan DM, Mossialos E: **A systematic review and critical assessment of incentive strategies for discovery and development of novel antibiotics.** *J Antibiot* 2016, **69**:73-88.
60. Czaplewski L, Bax R, Clokie M, Dawson D *et al.*: **Review alternatives to antibiotics—a pipeline portfolio review.** *Lancet Infect Dis* 2016, **16**:239-251.