Antimicrobial resistance three ways: healthcare crisis, major concepts, and the relevance of biofilms

Paula Jorge*, Andreia Patrícia Magalhães*, Tânia Grainha, Diana Alves, Ana Margarida Sousa, Susana Patrícia Lopes, Maria Olívia Pereira**

CEB - Centre of Biological Engineering, LIBRO - Laboratory of Research in Biofilms Rosário Oliveira, University of Minho, Campus de Gualtar, 4710-057, Braga, Portugal

* Paula Jorge and Andreia Patrícia Magalhães should be considered joint first authors
** Corresponding author

E-mail address: mopereira@deb.uminho.pt
Tel.: +351 253 604 402

Keywords
Antimicrobial Resistance, Tolerance, Persistence, Biofilms, Quorum Sensing, Polymicrobial Infection

Abstract
Worldwide, infections are resuming their role as highly effective killing diseases, as current treatments are failing to respond to the growing problem that is antimicrobial resistance (AMR). The social and economical burden of AMR seems ever rising, with health- and research-related organizations rushing to collaborate on a worldwide scale to find effective solutions. Resistant bacteria are spreading even in first-world nations, being found not only in healthcare-related settings, but also in food and in the environment. In this mini-review, the impact of AMR in healthcare systems and the major bacteria behind it are highlighted. Ecological aspects of AMR evolution and the complexity of its molecular mechanisms are explained. Major concepts, such as intrinsic, acquired, and adaptive resistance, as well as tolerance and heteroresistance, are also clarified. More importantly, the problematic of biofilms and their role in AMR, namely its main resistance and tolerance mechanisms, is elucidated. Finally, some of the most promising anti-biofilm strategies being investigated are reviewed. Much is still to be done regarding the study of AMR and the discovery of new anti-biofilm strategies. Gladly, considerable research on this topic is generated every day and increasingly concerted actions are being engaged globally to try and tackle this problem.
Introduction

Antibiotic discovery had an unprecedented role in medical advances, saving countless lives by mitigating infectious diseases, but the rapid global emergence of resistant bacteria over the last decades has been imperilling their worth (Martens and Demain 2017; WHO 2017a). Humankind is witnessing as antimicrobial resistance (AMR) becomes one of the biggest threats to medicine today, killing around 700 000 people worldwide each year (O’Neill 2014; Aslam et al. 2018). The aetiology of AMR is multifaceted, embracing (i) over consumption of antibiotics due to over prescription, self-medicating, or over-the-counter accessible antibiotics, (ii) absence of standardized guidelines for antibiotic usage, (iii) lack of sanitation/hygiene practices, and (iv) access to counterfeit drugs (Morgan et al. 2011; Laxminarayan and Heymann 2012; CDC 2013; Nature Editorial 2013; Luyt et al. 2014; Read and Woods 2014; Ventola 2015). Food is also an important source of AMR (Marshall and Levy 2011), due to the widespread use of antibiotics in animals, while the increased international human travelling and animal transportation aids in AMR spreading (EFSA 2018).

In healthcare settings, the concurrence of factors such as high antibiotic consumption, vulnerable patients, invasive practices, and inflow of pathogenic species have contributed for the substantial health and economic burden of AMR (Golkar, Bagasra and Pace 2014; Roca et al. 2015). To mitigate the increasing rate of AMR, main stakeholders (i.e. policy makers, public health authorities, regulatory agencies, pharmaceutical companies, and the scientific community) were prompted to take a concerted action. Therefore, measures, such as (i) rational/prudent use of antibiotics (Lushniak 2014), (ii) effective infection control measures, (iii) mitigation of environmental exposure, (iv) better diagnostic tools (Michael, Dominey-Howes and Labbate 2014), (v) prevention/surveillance research, and (vi) development of new therapies (Roca et al. 2015) were proposed.

Initiatives and programs raising awareness and promoting strategies to improve knowledge and reflections regarding AMR are key to fight its dissemination. Noteworthy, initiatives include the B-Debate (https://www.bdebatet.org), which fosters the dialogue amongst world-renowned multidisciplinary scientists on the growing threat of AMR at all health, animal, and environmental levels. In addition, the Joint Programming Initiative on Antimicrobial resistance (https://www.jpiamr.eu) has been defining a strategic research agenda under the assumption that only collaborative effort by an interdisciplinary team will afford the necessary critical mass and the scientific expertise to tackle AMR. Likewise, different agencies across the globe are engaged to make all efforts to control AMR. These agencies include the Global Antimicrobial Resistance Surveillance System (GLASS) (https://www.who.int/glass), the Centers for Disease Control and Prevention (CDC) (https://www.cdc.gov), the Food and Agriculture Organization (FAO) (http://www.fao.org), the European Centre for Disease Prevention and Control (ECDC) (https://ecdc.europa.eu/), the European Medicines Agency (EMA) (https://www.ema.europa.eu/), the World Alliance Against Antibiotic Resistance (WAAAR) (https://www.waaar.org/), the Global Health Security Agenda (GHSA) (https://www.ghsagenda.org/), and many others. Despite the proposed recommendations and resolutions, little progress has been made so far, and AMR shows no signs of decline.
**Multidrug-resistant organisms: the “superbugs”**

AMR evolution is nothing but Darwinian selection. Microorganisms evolved to develop mechanisms to escape lethal effects of antimicrobials (Forsberg *et al.* 2014; Aslam *et al.* 2018). Notably, the aberrant use of antibiotics exerted a significant selective pressure for the development of multidrug-resistant (MDR) organisms. These “superbugs” are able to resist multiple classes of antibiotics, evading the majority of current therapies (Stokes and Gillings 2011; Munita and Arias 2016) and spreading at an alarming rate, leading to abnormal rates of morbidity/mortality (Khameneh *et al.* 2016). A recent 2018 study shows that “superbug” infections accounted for 33 000 annual deaths in Europe in 2015, a burden that has been compared to that of other diseases combined (e.g. tuberculosis, HIV, flu) (Cassini *et al.* 2018).

MDR infections may be triggered by Gram-negative or -positive bacteria or even by fungal species. *Staphylococcus aureus* and Enterococcus species are among the most notorious “superbugs”, currently posing a pandemic threat (Watkins, David and Salata 2012; CDC 2013; Rossolini *et al.* 2014; Aslam *et al.* 2018). The most worrisome is the methicillin-resistant *S. aureus* (MRSA), whose ability to evolve and adapt to multiple settings (e.g. healthcare, community, livestock) has caused its rapid dispersal over the globe (Monaco *et al.* 2016). The spread of vancomycin-resistant enterococci (Golkar, Bagasra and Pace 2014) as well as the global epidemic of resistant *Streptococcus pneumoniae* and *Mycobacterium tuberculosis* (common respiratory pathogens) (Rossolini *et al.* 2014) also represent serious threats. Similarly, the emergence of MDR Gram-negative pathogens, typically thriving in healthcare facilities, namely Enterobacteriaceae (mostly *Klebsiella pneumoniae*), *Pseudomonas aeruginosa*, *Acinetobacter* spp., extended-spectrum β-lactamase (ESBL)-producing *Escherichia coli* and *Neisseria gonorrhoeae* (Rossolini *et al.* 2014), is particularly worrying (CDC 2013; Golkar, Bagasra and Pace 2014). In 2017, the World Health Organization (WHO) issued a global priority pathogens list (PPLs) of antibiotic-resistant bacteria (WHO 2017a) to help in prioritizing the research and development of new and effective antibiotic treatments, updating the previous PPLs issued by the CDC in 2013 (CDC 2013). Accordingly, the WHO has stratified the resistant pathogens in three priority tiers: “critical”, “high”, and “medium”. Table 1 summarizes the key features regarding these top bacterial threats.

**What is AMR?**

Understanding the evolution, divergence, and spread of AMR, along with the mechanisms behind it, is the main step in predicting and preventing this threat. In addition, it is important to understand the underlying concepts of AMR, such as resistance, heteroresistance, and tolerance, to facilitate knowledge dissemination and integration in the development of novel strategies to defeat it. Resistance, although tolerance may fit some of the criteria, is mainly classified in three forms: intrinsic, acquired, or adaptive. Frequently, microorganisms exhibit more than one form of resistance simultaneously, greatly contributing to the difficulty in finding suitable treatments. As such, all these aspects are discussed next.
Ecological evolution of AMR

Since the beginning of the antibiotic age, with the introduction of penicillin in the 1940s (Gaynes 2017), that researchers and physicians have been made aware of how strongly and quickly microorganisms fight back. Indeed, in 1941, penicillin was first administered and, in 1942, penicillin resistant bacteria were detected. Similarly, methicillin was introduced in 1960 and, in 1961, methicillin resistance was reported. With resistant strains propagating in this increasingly rapid pace, antibiotic utilization quickly led a golden era of Medicine to the current AMR crisis (Landecker 2016).

Regardless of its clear impact in modern medicine, however, AMR is actually an ancient and natural phenomenon, as microorganisms always had to defend themselves against naturally occurring antibiotics, with AMR evolving alongside their production (Perry, Westman and Wright 2014). In reality, several studies have revealed the existence of resistance genes in microorganisms preceding the antibiotic era. For instance, genes encoding resistance to natural antibiotics, namely β-lactams, tetracyclines, and glycopeptides, were found in 5000 and 30000-year-old permafrost sediments (D’Costa et al. 2011; Perron et al. 2015). Surprisingly, resistance genes against modern semi-synthetic antibiotics that do not occur naturally in microorganisms, namely amikacin, were also found (Perron et al. 2015). More recently, *Paenibacillus* sp. LC231 from a 4 Myr old isolated cave was found to harbour conserved resistance genes to most clinically used antibiotics (Pawlowski et al. 2016). These studies demonstrate that AMR is a natural phenomenon preceding the modern selective pressure of antibiotics, which may be simply selecting for pre-existing, hence intrinsic, determinants in the resistome (i.e. the resistance genetic pool of all microorganisms).

Besides its ancient intrinsicality, the major issue of AMR is its ability to spread from one microorganism to another. Although it was first believed that AMR was only inherited vertically within a resistant population, researchers quickly realized that bacteria were able to acquire resistant determinants from other bacteria by horizontal gene transfer, as further explained below. This ability to exchange genetic material has been the great source of bacterial genetic variation over time, in which the resistome acts as a widely available and sharable resource (Landecker 2016).

Despite AMR dissemination being primordial, its frequency and distribution has suffered an anthropogenic impact and changed historically, driven and sustained by the scale of antibiotic usage in clinical, veterinary, husbandry, and agricultural settings. For example, due to the large use of antibiotics, host microbiota, although harmless, nowadays carries a high content of resistance genes specific for the antibiotics used in medical and food production settings of a given country (Forslund et al. 2014). This creates a genetic pool that facilitates microbial pathogens to acquire resistance determinants when in contact with the host. The rate of bacterial release and uptake of genetic material and genetic recombination is also accelerated in the presence of an external stress like antibiotics, but also heavy metals and disinfectants coming from industrial settings (Landecker 2016).

Resistance genetic elements originating from anthropogenic sources are not only spreading within host microbiota, but also to the environment, including remote areas with minimal antibiotic exposure (Bartoloni et al. 2004, 2009; Pallecchi et al. 2008; McCann et al. 2019), very likely due to waste streams emanating from human activity. In truth, a great portion of antibiotics used for humans and animals are excreted and released unchanged into the environment, either due to
incomplete metabolization or due to disposal of unused drugs into the sewer, greatly contributing for the high load of antibiotics encountered in the environment today (Gillings and Stokes 2012).

Intrinsic AMR
Intrinsic AMR refers to the innate ability of microorganisms to resist to a specific antimicrobial due to genes in their genome encoding inherent structural or functional traits that provides them protection (Blair et al. 2015). This is evident in the disparate efficacies of most antibiotics against Gram-negative vs -positive bacteria due to their inherent distinct cell wall composition acting as barrier to the entrance of antibiotics into the cells (Arzanlou, Chai and Venter 2017; Petchiappan and Chatterji 2017). For instance, vancomycin, a common antibiotic in MRSA treatment, is effective against Gram-positives because it easily reaches their peptidoglycan cell wall. Due to constrains in overcoming the outer membrane of Gram-negatives, vancomycin is ineffective against these bacteria (Rice 2012). Similarly, daptomycin is active against Gram-positives, but it is unable to act against Gram-negative bacteria due to the lower proportion of anionic phospholipids in their cytoplasmic membrane (Randall et al. 2013). Several studies are tackling this issue by modifying existing compounds to make them active against Gram-negative bacteria. For example, a recent study was successful in optimizing arylomycins, a weak class of natural products, to produce a potent, broad-spectrum molecule, G0775, active against Gram-negative bacteria by inhibiting the essential bacterial type I signal peptidase (Smith et al. 2018). Another example is the conversion of the natural product deoxyxybomycin, only active against Gram-positives, into an antibiotic able to accumulate inside and be active against Gram-negatives, with the help of computational modelling (Richter et al. 2017).

High levels of AMR can be achieved through intrinsic restricted or selective outer membrane permeability, drug efflux pump systems, and/or expression of intrinsic antibiotic resistance genes (Blair et al. 2015). Bacteria can limit the entry of broad-spectrum drugs, e.g. carbapenems and cephalosporins, by reducing or replacing non-specific porin proteins by specific or more selective protein channels (Nikaido 2003; Fernandez and Hancock 2012). For instance, P. aeruginosa deficient in OprD porin, responsible for diffusion of small peptides, displays resistance to carbapenems (Pechère and Köhler 1999). This kind of bacterial mechanism is well studied and it has been reviewed previously (Kumar and Schweizer 2005; Langton, Henderson and Herbert 2005; Poole 2005). Drug efflux systems are protein complexes located in the cell wall of Gram-negative bacteria responsible for expelling toxic molecules as antibiotics. Several bacteria possess genes encoding efflux pumps, greatly contributing to AMR (Blair, Richmond and Piddock 2014; Sanchez-Romero and Casadesus 2014). P. aeruginosa is a well-characterized example with clinically relevant efflux pumps such as the MexAB-OprM and MexXY/OprM(OprA) systems, which contribute to a stable and consistent resistance to a wide range of antibiotics and protection against molecules targeting the ribosomal machinery, respectively (Li et al. 1994; Li, Livermore and Nikaido 1994; Li, Nikaido and Poole 1995).

Microorganisms can also be intrinsically resistant to antibiotics due to the expression of antibiotic resistance genes (Liu et al. 2010; Blake and O’Neill 2013; Xu et al. 2017; Peterson and Kaur 2018). For instance, β-lactam antibiotics have no action against M. tuberculosis because these bacteria inherently produce β-lactamases, such as BlaC, that hydrolyse the β-lactam ring inactivating this class of antibiotics (Smith, Wolff and Nguyen 2012). Another example of intrinsic resistance is the absence of a susceptible target site for an antibiotic to act on (Blair et al. 2015). For instance, the
Biocide triclosan is ineffective against *P. aeruginosa* because it carries the *fabV* gene encoding a triclosan-resistant enoyl-ACP reductase, the site of action of triclosan (Zhu *et al.* 2010).

**Acquired AMR**

Microorganisms can acquire or develop resistance, being this what most greatly contributes to the AMR crisis (Blair *et al.* 2015). Acquired resistance arises when an originally antibiotic-sensitive organism becomes resistant through the acquisition and incorporation of new genetic material (e.g. plasmids, transposons, integrons, naked DNA) from other microorganisms by horizontal gene transfer or as a result of mutations of chromosomal (intrinsic) genes (Arzanlou, Chai and Venter 2017; Pang *et al.* 2019). The spread of β-lactam resistance among bacteria is the major example, as several species are able to acquire plasmids encoding β-lactamase genes leading to the emergence of, for e.g., ESBL- and metallo-β-lactamase-producing *P. aeruginosa*, ESBL-producing *E. coli*, *Haemophilus influenzae*, *N. gonorrhoeae*, *Salmonella*, *Shigella*, and *Vibrio cholera* (Rawat and Nair 2010).

In general, acquired resistance can be mediated by (i) reduced antibiotic uptake and (ii) increased antibiotic efflux, reducing its intracellular concentration, (iii) antibiotic modification or inactivation, and (iv) antibiotic target modification by genetic mutation or post-translational modification. Often, these mechanisms are combined, contributing to the expression of high levels of AMR, as is the case of increased resistance observed against β-lactams (Arzanlou, Chai and Venter 2017). As intrinsic resistance, acquired resistance is stable and transmitted vertically (Blair *et al.* 2015). This vast topic is only outlined here, but is reviewed in detail in (Nikaido 2009; Blair *et al.* 2015).

Frequently, acquired and intrinsic mechanisms are closely related, as mutations can alter the expression of intrinsic resistance mechanisms. For instance, carbapenem resistance in Enterobacteriaceae generally involves the production of β-lactamases. Nevertheless, if mutations in porin production occur, bacteria can reduce or even end β-lactamase production (Wozniak *et al.* 2012; Lavigne *et al.* 2013; Tangden *et al.* 2013). Another example is that mutations can enhance *P. aeruginosa* intrinsic antibiotic resistance through loss of oprD porin expression, via mutation in the *oprD* gene or its associated regulatory proteins, and de-repression of chromosomal AmpC β-lactamase and MexAB-OprM multidrug efflux pump, conferring resistance to β-lactam antibiotics (Taylor, Yeung and Hancock 2014). Efflux pumps are one of the major contributors to intrinsic resistance that microorganisms can mobilize onto plasmids and transfer to other bacteria. For instance, IncH1 plasmid isolated from *Citrobacter freundii* includes genes encoding the New Delhi metallo-β-lactamase 1, but also a tripartite resistance nodulation division pump (Dolejska *et al.* 2013).

**Adaptive AMR**

Adaptive AMR is one of the most complex forms of bacterial resistance (Fernández, Breidenstein and Hancock 2011). It consists in the ability to alter gene or protein expression very rapidly in response to an antibiotic insult or environmental cues, such as pH, temperature, nutrient, or oxygen limitation (Fernández, Breidenstein and Hancock 2011; Motta, Cluzel and Aldana 2015; Arzanlou, Chai and Venter 2017). Development of adaptive AMR in the presence of antibiotics is usually observed when cells are exposed first to non-lethal levels of such agents, but may escalate to where bacteria are able to survive otherwise lethal concentrations if they are consecutively exposed to increasing antibiotic doses. In fact, bacteria can increase their level of resistance gradually and across...
generations if the stimulus endures, indicating the existence of some type of resistance memory (Sandoval-Motta and Aldana 2016).

Unlike intrinsic and acquired resistance, adaptive resistance is unstable, transient, highly dependent of the presence of antibiotics, and it cannot be vertically transmitted. After removal of the triggering factor, microorganisms revert to their "original state", re-gaining susceptibility, although the original level of resistance may not be restored (Fernández, Breidenstein and Hancock 2011; Arzanlou, Chai and Venter 2017; Pang et al. 2019). Because of this, adaptive AMR has been linked with the phenomenon of MIC baseline creep seen in many bacterial species, in which the average minimum inhibitory concentration (MIC) increases from the moment of antibiotic introduction onward, making them more likely to achieve the resistance breakpoint over time (Fernández, Breidenstein and Hancock 2011).

Because of its transient nature, adaptive resistance represents one of the biggest challenges in designing effective antimicrobial therapies, explaining the common differences found between in vitro and in vivo antibiotic susceptibilities (Fernández, Breidenstein and Hancock 2011). Adaptive resistance represents a crucial biological advantage and an intelligent survival mechanism since microorganisms do not pay the fitness costs associated with irreversible mutations (Motta, Cluzel and Aldana 2015), reverting to their “original state” when more advantageous (Andersson and Hughes 2010; Motta, Cluzel and Aldana 2015).

There are several mechanisms of adaptive resistance, including epigenetic inheritance, population heterogeneity, mutability, gene amplification, efflux pumps, and biofilm formation (Sanchez-Romero and Casadesus 2014; Motta, Cluzel and Aldana 2015). The molecular mechanisms behind adaptive resistance are still poorly understood but apparently quite complex, involving intricate regulatory pathways. Moreover, adaptive resistance may interplay with intrinsic or acquired resistance (Fernández, Breidenstein and Hancock 2011) as the genetic mutations or epigenetic changes triggered by environmental conditions can influence the expression of intrinsic and acquired mechanisms of resistance (Sanchez-Romero and Casadesus 2014; Motta, Cluzel and Aldana 2015). A great example of this phenomenon is shown in a recent study, where a sub-population of E. coli cells with increased expression of efflux pumps were found to also have a higher mutability rate due to the decrease expression of a the DNA mismatch repair gene, which led to mutants with higher antibiotic resistance (El Meouche and Dunlop 2018). Biofilm formation is a “perfect” example of the interplay of the three types of resistance. Bacteria undergo genetic and phenotypic alterations to adhere and produce an exopolymeric matrix to bind to a surface and to other bacteria (Stewart 2014; Donné and Dewilde 2015; Kumar et al. 2017).

**Resistance, heteroresistance, and tolerance**

Despite very commonly encountered in AMR studies, the concepts of resistance, heteroresistance, and tolerance are sometimes misused, being of importance to elucidate them. Resistance refers to the ability of microorganisms to survive and grow at increased antibiotic concentrations for long periods and it is quantifiable by assessing the MIC (Brauner et al. 2016). However, it sometimes happens that different antibiotic susceptibilities exist within the same bacterial population, which can include resistant subpopulations. This phenomenon is termed heteroresistance and, although generally disregarded in clinical settings, it is critical in foreseeing the success of a given antimicrobial therapy, since poor designed treatments may select for the resistant populations.
Heteroresistance is usually detected in MBC, disc diffusion, or e-test assays when discrete colonies grown in the zone of inhibition, and can be confirmed by a population analysis profiling assay (El-Halfawy and Valvano 2015).

In turn, tolerance refers to the ability of microorganisms to survive a transient exposure to increased antibiotic concentrations, even those above the MIC. However, unlike resistance, tolerance is only temporary, as it just takes more time for the antibiotic to kill bacteria. Tolerance can be due to slow growth, which in turn can be inherent, i.e. characteristic of a given species or strain, or non-inherent, i.e. caused by poor growth conditions (e.g. biofilms), stress factors (e.g. antibiotics), or by bacterial stationary growth phase. Tolerance, however, may also be due to antibiotic application to a bacterial population in the lag growth phase, in which they are transitioning from growth-arrest to an exponential growth phase (Brauner et al. 2016; Balaban et al. 2019). For more detailed information about resistance and tolerance definitions, the two cited reviews are recommended. In the next section, focus will be given to biofilms and their resistance and tolerance traits.

Biofilm resistance and tolerance

What are biofilms?
Contrary to the typical idea of single-species free-floating bacteria, microorganisms naturally reside in groups, establishing complex and dynamic consortia called biofilms. The ability of microorganisms to persist and thrive within biofilms is an important feature denoting critical concern in clinical settings. Indeed, biofilms play a significant role in microbial survival and persistence in natural ecosystems, thus being ubiquitous in Nature and considered an ancient form of microbial adaptation. Remarkably, it is speculated that the transition of microorganisms to the biofilm mode of growth established the first multicellular life form as an adaptive response to the extreme conditions encountered in early Earth (de la Fuente-Núñez et al. 2013).

Biofilms are usually characterized as well-organized structures of microorganisms attached to biotic or abiotic surfaces and whose cells are encased and protected by a self-produced polymeric matrix. Typically, the biofilm life cycle encompasses three stages, namely (i) attachment, (ii) maturation, and (iii) dispersion. The first stage initiates by the reversible binding of bacteria to a surface followed by their irreversible attachment. Next, bacterial growth and matrix production take place, leading to increased biomass and maturation of the biofilm. At this stage, biofilms develop microenvironments dependent on nutrient and oxygen gradients, with cells developing different phenotypes depending on their spatial organization. Finally, biofilms eventually disperse, allowing cells to migrate and colonize other areas (Bjarnsholt et al. 2013).

Concerning the impact of biofilms in healthcare settings, it is important to realize that the vast majority infections are actually biofilm-mediated (Høiby et al. 2015). Biofilm infections can be device-related (e.g. catheters, implants, contact lenses, prosthetic valves and joints) or tissue-related (e.g. endocarditis, chronic otitis media, lung infections in cystic fibrosis, chronic wounds) (Lebeaux, Ghigo and Beloin 2014). In these infections, the physiological features of biofilm cells and the matrix surrounding them contribute to their higher resistance/tolerance to external stresses, including the action of antimicrobials and the immune system, allowing the establishment of persistent/chronic infections (Grant and Hung 2013; Donné and Dewilde 2015; Kumar et al. 2017). Moreover, most
Biofilm infections normally have a polymicrobial aetiology, with phylogenetically different microorganisms co-existing (Peters et al. 2012; Giaouris et al. 2015; Costa-Orlandi et al. 2017). The polymicrobial nature of most biofilm-mediated infections can lead to the chronic scenario of infection (Stacy et al. 2016), as the interactions amongst the resident species may augment the severity of the infection and contribute for the recalcitrance towards conventional therapy (Wolcott et al. 2013; Schroeder, Brooks and Brooks 2017). Biofilms employ different yet concerted resistance and tolerance mechanisms illustrated in Figure 1 and further detailed in the next sections.

Extracellular matrix

Microorganisms living in a biofilm are surrounded and encased by a majorly self-produced matrix, which can comprise over 90% of the total mass of the biofilm (Flemming and Wingender 2010). The biofilm matrix is a complex and intricate amalgamation of different hydrated extracellular polymeric substances (EPS), including polysaccharides, proteins, nucleic acids, and lipids. These molecules offer biofilms their structure and mechanical stability by forming a three-dimensional network that supports biofilm adherence and cell immobilization (Flemming and Wingender 2010). The matrix and its constituents make up the first barrier to the entry and diffusion of foreign substances into the biofilm, often impeding them from reaching the cells, thus greatly prompting biofilm antimicrobial resistance (Figure 1). Yet, and despite its significance, antimicrobial penetration hindrance does not fully explain the resistance phenomena seen in biofilm scenarios, with some antibiotics rapidly reaching the biofilm cells while failing at compromising their viability (Hall & Mah, 2017). As explained in the next sections, the mechanisms through which resistance and tolerance appear in biofilms is complex.

A major and important matrix component is extracellular DNA (eDNA), ubiquitous to almost all biofilms and with structural and cell-to-cell interconnecting functions (Whitchurch et al. 2002; Barken et al. 2008). eDNA has shown to confer protection from aminoglycosides to P. aeruginosa biofilms, most likely due to its electrostatic interaction with positively charged antibiotics (Chiang et al. 2013). Notably, the presence of an antimicrobial can itself promote the eDNA release by the biofilm cells to the matrix. For instance, biofilms of Staphylococcus epidermidis doubled their amount in eDNA when treated with vancomycin, thus benefiting from its affinity for the positively charged antibiotic, which was prevented from reaching the cells and exerting its activity (Doroshenko et al. 2014). eDNA has also been shown to induce the expression of resistance genes. This occurs by chelating cations such as Mg$^{2+}$ and by creating an acidic micro domain around itself, two environmental signals that activate signalling pathways linked to antimicrobial resistance, such as the PhoPQ and PmrAB two component systems in S. Typhimurium (Johnson et al. 2013) and P. aeruginosa (Wilton et al. 2016). Furthermore, eDNA has also been related with increased horizontal gene transfer in biofilms, serving as vehicle for resistance genes and causing the rapid spread of resistance between competent biofilm cells (Okshevsky and Meyer 2015).

Other matrix components affecting biofilm resistance are polysaccharides, crucial matrix components influencing biofilm adhesion and structure while also offering protection against antimicrobials. For example, the polysaccharide Psl from P. aeruginosa has shown to provide resistance to colistin, polymyxin B, tobramycin, and ciprofloxacin probably via electrostatic interactions, and this protective effect was extended to non Psl-producing species, such as E. coli and S. aureus (Billing et al. 2013). The biofilm matrix can also contain secreted antibiotic-modifying enzymes. For instance, secreted β-lactamases were able to degrade the antibiotic ampicillin in K.
*pneumoniae* biofilms, impeding it from reaching the cells in the biofilm (Anderl, Franklin and Stewart 2000).

**Nutritional constraints and persister cells**

Biofilms are a complex architectural conglomerate, not only due to their diverse composition in terms of EPS but also for possessing heterogeneous microhabitats caused by the establishment of an oxygen and nutrient gradient. This gradient derives from the faster oxygen and nutrient consumption compared to their diffusion rates through the biofilm, causing biofilm cells to appear stratified according to oxygen and nutrient availability. Remarkably, oxygen and nutrient depletion in lower layers can cause biofilm cells to adopt a low metabolic state or even cause cell death (Flemming *et al.* 2016). This gradient phenomenon partially explains the physiological heterogeneity encountered in most biofilms, characterised by cells with diverse gene expressions, metabolic activities and phenotypes, which translates into different AMR and tolerance abilities.

A great example of how nutritional constraints affect biofilm tolerance to antimicrobials is the case of cells that reduce their metabolic activity and enter a slow growth or dormancy state when nutrients and oxygen are low or absent, achieving the phenotype of persisters (Hall and Mah 2017). This type of persistence is known as time-dependent persistence or “tolerance by slow growth” (Brauner *et al.* 2016). Typically, most antimicrobials act best on fast-growing metabolically active cells. For example, β-lactams act by preventing the reassembly of the peptidoglycan layer bonds during bacterial growth, causing cell lysis (Horne and Tomasz 1977), whilst fluoroquinolones inhibit DNA gyrase, causing DNA damage (Crumplin and Smith 1976). Persisters are able to diminish the antimicrobial effectiveness of these and other antimicrobials without any genetic changes by simply stopping their metabolism and growth (Olsen 2015).

Persistence is characterized by occurring only in a subset of cells that usually comprise less than 1% of the microbial population of a biofilm, with antimicrobials only effectively killing the remainder of the biofilm cells (Brauner *et al.* 2016). As such, the persister phenotype is a major reason why certain antimicrobials are ineffective despite being able to reach the cells within a biofilm and is one of the main contributors to biofilm infection relapsing, as the surviving cells can regrow after antimicrobial treatment and maintain the infection (Conlon 2014). Gladly, efforts are being made to target these specific and troublesome bacterial subpopulations. For example, a recent study was successful in achieving total persister eradication by activating the ClpP protease with the acyldepsipeptide antibiotic (ADEP4). This made the protease non-specific, leading to persister cells self-digestion. Furthermore, ADEP4 combination with rifampicin was able to completely eradicate an *in-vivo* *S. aureus* biofilm infection (Conlon *et al.* 2013).

Unlike resistant bacterial populations, persistence is characterized by a biphasic killing curve, which translates the different rates that bacteria are killed within the same population. Also, persistent bacteria, unlike resistant ones, are unable to replicate in the presence of an antimicrobial, a characteristic that also differentiates this phenomenon from the one of heteroresistance (Balaban *et al.* 2019). Despite its importance, the molecular mechanisms behind the changes from susceptible to persister phenotypes are still being unravelled. Persisters have also been linked to ATP depletion in *S. aureus* (Conlon *et al.* 2016), *E. coli* (Shan *et al.* 2017), and *P. aeruginosa* (Cameron *et al.* 2018). A more recent study showed that the *msaABCR* operon, previously linked to virulence, biofilm development and antibiotic resistance, is involved as well (Sahukhal, Pandey and Elasri 2017).
Another recent finding shows that the presence of antimicrobials can induce the persister phenotype, namely through the putative de-N-acetylase DnpA after *P. aeruginosa* biofilm exposure to fluoroquinolones (Khandekar et al. 2018).

Interestingly, it has been recently described the occurrence of persister cell memory, by which cells of *E. coli*, *Acinetobacter radioresistens*, *S. Typhimurium*, *S. epidermidis* and *Bacillus subtilis* retain their persister phenotype up to weeks after being removed from biofilm cultures (Miyaue et al. 2018). Persister are characterized by presenting temporary thus reversible tolerance towards antimicrobial treatment, but the length in which cells remain in a persistent state due to the described memory effect gives them an extra advantage for surviving in antimicrobial-containing environments. For more detailed information about persister cells, the reviews by Conlon, Rowe and Lewis 2015, Brauner et al., 2016, Van den Bergh et al., 2017, Fisher et al., 2017, and Balaban et al. 2019 are recommended.

In addition to causing the described phenotypic changes in biofilm cells, low availability of oxygen, or hypoxia, has been also related to the expression of resistance related genes. Specifically, the *mexEF-oprN* and *mexCD-oprJ* efflux pump genes are upregulated in *P. aeruginosa* in low oxygen conditions (Schaible, Taylor and Schaffer 2012; Tata et al. 2016). Additionally, hypoxia may further protect biofilms from antimicrobials by impairing the formation of reactive oxygen species (ROS), which have been linked to cell killing by bactericidal antibiotics (Hall and Mah 2017). Low nutrient concentration, specifically amino acids, may also enhance biofilm tolerance through the stringent response, in which an accumulation of uncharged tRNAs triggers the production of the alarmone stress signal guanosine tetraphosphate (ppGpp). This response causes an induction of a lag phase (transient growth arrest), which has been showed to improve tolerance to antibiotics (Brauner et al. 2016; Hall and Mah 2017). Of notice, a recent study as show that low pH, another environmental factor, has been proven to cause the latter effect (Vulin et al. 2018).

### Quorum sensing

The mechanisms by which microorganisms within a biofilm regulate their activities are coordinated through a cell-to-cell communication system known as quorum-sensing (QS). QS is used by bacteria (and fungi) to sense population density and regulate gene expression accordingly, serving as channel for intra- and inter-species communication, allowing the establishment of intimate relationships of competition or cooperation, but also for communication between the microorganisms and its host (Li and Tian 2012; Grandclément et al. 2016). Microorganisms regulate these activities by releasing, sensing, and responding to small QS signal molecules termed auto-inducers (AI). When AI concentration reaches a threshold due to an increase in population density, these signal molecules activate receptors with the ability to alter gene expression, promoting beneficial behaviours under a particular condition, such as virulence factor expression, motility, and biofilm formation (Grandclément et al. 2016; Hawver, Jung and Ng 2016; Knecht et al. 2016). Using QS, microorganisms can switch from acting as individual cells to operating in a concerted multicellular fashion, thereby switching to the biofilm mode of growth and accommodating to or escaping from antimicrobial stresses (Filkins and O’Toole 2015; Passos da Silva et al. 2017).

QS plays a key role in biofilm AMR, although the mechanisms behind it are still being unravelled. QS influences the production of EPS, which are key players in biofilm resistance, as described above. For example, the PqsABCDE/PqsR QS system in *P. aeruginosa* stimulates the production of eDNA (Pérez-
Pérez et al. 2017), which is highly related with AMR in biofilms, as described previously. QS as also been linked to the upregulation of resistance genes, as is the case of oxacillinase S1, AmpC, AdeA and AdeB in Acinetobacter baumannii (Dou et al. 2017). More recently, Chromobacterium violaceum was reported to use QS to increase its resistance to bactobolin, a Burkholderia thailandensis antibiotic, by increasing transcription of a putative antibiotic efflux pump (Evans et al. 2018).

As stated, QS may serve inter-species communication, with the AI from one species interfering with signalling pathways of other species present in the same biofilm, thus altering gene expression or directly affecting the physiology of the co-habitants (Schertzer, Boulette and Whiteley 2009; Elias and Banin 2012). The role of inter-species communication in biofilm resistance is further explored next.

**Inter-species interactions**

Most of the research on biofilm resistance has been focused on single-species biofilms. However, these simple laboratory models do not illustrate the true nature of biofilm communities, since most of biofilm-mediated infections are actually polymicrobial (Wolcott et al. 2013; Gabrielska and Rumbaugh 2015). The inclusion of the multispecies factor in AMR studies is pivotal, as it is becoming increasingly clear that interactions between different species can modulate the overall consortium behaviour, resulting in enhanced AMR and infection severity (Dalton et al. 2011; Peters et al. 2012; Murray et al. 2014; Bowen et al. 2018).

By enclosing multiple species, biofilms obtain numerous ecological advantages, with established interactions, either cooperative or competitive, usually resulting in a beneficial outcome to the biofilm. A cooperative interaction is, for example, the metabolite cross-feeding between Aggregatibacter actinomycetemcomitans and Streptococcus gordonii that benefits the latter while also enhancing A. actinomycetemcomitans virulence (Ramsey, Rumbaugh and Whiteley 2011).

Regarding competitive interactions, a great example is the one established between P. aeruginosa and S. aureus. P. aeruginosa produces the enzyme LasA that selectively lysis S. aureus, whose content in iron is used for P. aeruginosa growth, increasing its pathogenic potential under low-iron conditions (Mashburn et al. 2005). However, S. aureus growth is not completely inhibited by P. aeruginosa, with the latter inducing expression of virulence factors and facilitating the emergence of small colony variants in S. aureus (Mitchell et al. 2010). This phenotype allows S. aureus to survive in proximity with P. aeruginosa, being linked to infection persistence, establishment of intracellular infections, and lower antimicrobial susceptibility due to their reduced metabolic state (Garcia et al. 2013; Proctor et al. 2014). These interactions raise a healthcare concern, as polymicrobial biofilm infections are typically more severe and recalcitrant to treatment (Wolcott et al. 2013). For instance, P. aeruginosa and S. aureus co-infection delayed wound healing and triggered host inflammatory responses (Pastar et al. 2013). Also, P. aeruginosa displayed higher virulence when grown with Gram-positive bacteria in vivo (Korgaonkar et al. 2013).

Regarding specifically AMR, several works have emphasized the increasing resistance to antibiotics in multispecies biofilms (Adam, Baillie and Douglas 2002; Lopes et al. 2012; Lee et al. 2014; Magalhães, Lopes and Pereira 2017) and some examples are presented in Table 2. The studies reviewed suggest that mechanisms, such as interspecies signalling, biofilm matrix production, and horizontal gene transfer are major contributors to the increased multispecies biofilm resistance.
Since much is still unknown, it is imperative to continue the study of interspecies interactions (from a molecular standpoint) that lead to the increased AMR of biofilms.

**Anti-biofilm strategies**

A comprehensive knowledge of AMR mechanisms is crucial to find suitable anti-biofilm strategies. So far, these essentially belong to three different approaches, i.e. inhibition of bacterial attachment to surfaces, interference with signal molecules that modulate biofilm formation, and disruption of the biofilm architecture (Parrino *et al.* 2019), for which examples are given in Table 3.

As bacterial adhesion is the first step in biofilm formation, a number of surface modifications have been developed to prevent bacterial attachment and/or kill adhered bacteria through the immobilization of antimicrobials (Desrousseaux *et al.* 2013; Hasan, Crawford and Ivanova 2013; Alves *et al.* 2016). Among the antimicrobial agents explored in surface functionalization, antimicrobial peptides, enzymes, bacteriophages, and essential oils stand out as promising alternatives to antibiotics. These natural bactericidal compounds are considered to have a low propensity for the development of AMR due to their mechanisms of action (Glinel *et al.* 2012). For instance, antimicrobial peptides target the bacterial membrane, so their activity does not require cells to be metabolically active (Kumar, Kizhakkedathu and Straus 2018), allowing them to effectively kill cells that are dormant or non-growing, such as persister cells (Batoni, Maisetta and Esin 2016; Grassi *et al.* 2017). In turn, bacteriophages are natural bacterial predators, comprising a promising option to overcome biofilm barriers when used in combined therapies or after being genetically engineered with new functions to overcome biofilm obstacles (Pires *et al.* 2017).

Another anti-biofilm strategy is the interference with signal molecules that modulate biofilm development, where QS along with intracellular signalling by bis-(3'→5')-cyclic-dimeric guanosine monophosphate (c-di-GMP) have been the subject of great attention (LaSarre and Federle 2013; Parrino *et al.* 2019). QS interference can be achieved by degrading AI or inhibiting their production, limiting the activity of QS signal receptors, or mimicking AI using structurally synthetic compounds. A number of compounds targeting QS using these approaches have been identified, such as quorum sensing inhibitors that block the action of AI and quorum quenching enzymes that degrade signalling molecules (Hirakawa and Tomita 2013; Rémy *et al.* 2018; Kalia *et al.* 2019). It is postulated that resistance to these anti-QS strategies would develop slowly, making them promising alternatives to traditional antibiotics (Turkina and Vikström 2018). In turn, c-di-GMP has been described as a key mediator of biofilm formation, especially in Gram-negative bacteria, by stimulating the biosynthesis of adhesins, adhesive pili, and EPS, and by inhibiting bacterial motility. Since c-di-GMP is not essential for bacterial growth, its inhibition should not promote resistance, therefore being a good target for the development of anti-biofilm compounds (Valentini and Filloux 2016).

After bacterial attachment to a surface, large amounts of EPS are produced. Given its role in biofilm resistance, as described above, a promising strategy to dismantle established biofilms relies on the use of enzymes targeting EPS (Alves and Pereira 2014). Matrix disruptive enzymes, such as alginate lyase (Ramsey and Wozniak 2005), DNase I (Sugimoto *et al.* 2018), lysozyme (Ragland and Criss 2017), dispersin B (Kaplan *et al.* 2003), and lysostaphin (Bastos, Coutinho and Coelho 2010) have been extensively investigated with this aim, many times in combination with other antimicrobials.
Another strategy to target established biofilms relies on the stimulation of the natural stage of biofilm dispersal. Although this comprises a survival strategy of biofilms to colonize new areas, the dispersed and free cells are technically more susceptible to antimicrobials and host defences (Kostakioti, Hadjifrangiskou and Hultgren 2013). For instance, it has been demonstrated that a low concentration of c-di-GMP leads to biofilm self-destruction. As such, c-di-GMP should be considered as a good target for a biofilm dispersion strategy (O’Toole and Ha 2015). Despite promising results, most of these strategies fail to be tested and validated using *in vivo* models, so the development of strategies to fight biofilms are still urgently needed (Magana *et al.* 2018).

**Conclusion**

The solution for the hitches caused by one of the smallest life forms on this planet, i.e. bacteria, remains an unsolved riddle, as these microorganisms do not cease to amaze with their ability to circumvent every “curve ball” thrown their way. Their capability to evolve and adapt has led to a modern healthcare crisis as they become resistant to most, and sometimes all, available antibiotics. The complex issue of AMR, it seems, is a “many-fronts battle”, with biofilm formation being a substantial portion of the problem. This ancient form of bacterial adaptation is itself a form of AMR that escalates when resistant bacteria are its originators, making antibiotics forced to not only surpass bacterial resistance mechanisms (e.g. efflux pumps, cell-wall modifications) but also biofilm specific constrains (e.g. EPS matrix, persister cells).

The engagement from health-and research-related organizations worldwide is being put to the test, with many believing that only a concerted global action will result in AMR mitigation. In order to do so, unravelling the molecular mechanisms behind this phenomenon is pivotal in order to elaborate new effective antimicrobial strategies. Although much has yet to be done, substantial research is created every day to tackle this problem. Innovative solutions, such as surface functionalization to prevent biofilm formation, discovery of compounds that interfere with bacterial communication, and enzyme application to disperse grown biofilms are just a few examples. Slowly but surely we will come to a solution; let us hope it is not too late!

**Funding**

This work was supported by the Portuguese Foundation for Science and Technology (FCT) under the scope of the strategic funding of UID/BIO/04469/2019 unit and COMPETE 2020 (POCI-01-0145-FEDER-006684) and BioTecNorte operation (NORTE-01-0145-FEDER-000004) funded by the European Regional Development Fund under the scope of Norte2020 - Programa Operacional Regional do Norte. The authors also acknowledge COMPETE2020 and FCT for the project POCI-01-0145-FEDER-029841, and FCT for the PhD Grants of Andreia Magalhães [grant number SFRH/BD/132165/2017] and Tânia Grainha [grant number SFRH/BD/136544/2018].

**Conflicts of interest**

This work presents no conflicts of interest.
References

Adam B, Baillie GS, Douglas LJ. Mixed species biofilms of Candida albicans and Staphylococcus epidermidis. J Med Microbiol 2002;51:344–9.

Agnihotri S, Mukherji S, Mukherji S. Immobilized silver nanoparticles enhance contact killing and show highest efficacy: Elucidation of the mechanism of bactericidal action of silver. Nanoscale 2013;5:7328–40.

Alba C, Blanco A, Alarcón T. Antibiotic resistance in Helicobacter pylori. Curr Opin Infect Dis 2017;30:489–97.

Alves D, Magalhães A, Grzywacz D et al. Co-immobilization of Palm and DNase I for the development of an effective anti-infective coating for catheter surfaces. Acta Biomater 2016;44:313–22.

Alves D, Pereira MO. Mini-review: Antimicrobial peptides and enzymes as promising candidates to functionalize biomaterial surfaces. Biofouling 2014;30:483–99.

Anderl JN, Franklin MJ, Stewart PS. Role of antibiotic penetration limitation in Klebsiella pneumoniae biofilm resistance to ampicillin and ciprofloxacin. Antimicrob Agents Chemother 2000;44:1818–24.

Andersson DI, Hughes D. Antibiotic resistance and its cost: Is it possible to reverse resistance? Nat Rev Microbiol 2010;8:260–71.

Antonic V, Stojadinovic A, Zhang B et al. Pseudomonas aeruginosa induces pigment production and enhances virulence in a white phenotypic variant of Staphylococcus aureus. Infect Drug Resist 2013;6:175–86.

Arias CA, Murray BE. The rise of the Enterococcus: beyond vancomycin resistance. Nat Rev Microbiol 2012;10:266–78.

Arizpe A, Reveles KR, Patel SD et al. Updates in the management of cephalosporin-resistant Gram-negative bacteria. Curr Infect Dis Rep 2016;18:39.

Armbruster CE, Hong W, Pang B et al. Indirect pathogenicity of Haemophilus influenzae and Moraxella catarrhalis in polymicrobial otitis media occurs via interspecies quorum signaling. MBio 2010;1:e00102-10.

Arzanlou M, Chai WC, Venter H. Intrinsic, adaptive and acquired antimicrobial resistance in Gram-negative bacteria. Essays Biochem 2017;61:49–59.

Aslam B, Wang W, Arshad MI et al. Antibiotic resistance: a rundown of a global crisis. Infect Drug Resist 2018;11:1645–58.

Bachovchin DA, Brown SJ, Rosen H et al. Identification of selective inhibitors of uncharacterized enzymes by high-throughput screening with fluorescent activity-based probes. Nat Biotechnol 2009;27:387–94.

Balaban NQ, Helaine S, Lewis K et al. Definitions and guidelines for research on antibiotic
persistence. Nat Rev Microbiol 2019;17:441–8.

Banin E, Brady KM, Greenberg EP. Chelator-induced dispersal and killing of Pseudomonas aeruginosa cells in a biofilm. Appl Environ Microbiol 2006;72:2064–9.

Banin E, Vasil ML, Greenberg EP. Iron and Pseudomonas aeruginosa biofilm formation. PNAS 2005;102:11076–81.

Barken KB, Pamp SJ, Yang L et al. Roles of type IV pili, flagellum-mediated motility and extracellular DNA in the formation of mature multicellular structures in Pseudomonas aeruginosa biofilms. Environ Microbiol 2008;10:2331–43.

Barraud N, Storey M V., Moore ZP et al. Nitric oxide-mediated dispersal in single- and multi-species biofilms of clinically and industrially relevant microorganisms. Microb Biotechnol 2009;2:370–8.

Bartoloni A, Bartalesi F, Mantella A et al. High prevalence of acquired antimicrobial resistance unrelated to heavy antimicrobial consumption. J Infect Dis 2004;189:1291–4.

Bartoloni A, Pallecchi L, Rodríguez H et al. Antibiotic resistance in a very remote Amazonas community. Int J Antimicrob Agents 2009;33:125–9.

Bastos M do C de F, Coutinho BG, Coelho MLV. Lysostaphin: A staphylococcal bacteriolysin with potential clinical applications. Pharmaceuticals 2010;3:1139–61.

Boni G, Maisetta G, Esin S. Antimicrobial peptides and their interaction with biofilms of medically relevant bacteria. Biochim Biophys Acta - Biomembr 2016;1858:1044–60.

Van den Bergh B, Fauvart M, Michiels J. Formation, physiology, ecology, evolution and clinical importance of bacterial persisters. FEMS Microbiol Rev 2017;41:219–51.

Billings N, Millan M, Caldara M et al. The extracellular matrix Component Psl provides fast-acting antibiotic defense in Pseudomonas aeruginosa biofilms. PLoS Pathog 2013;9:e1003526.

Bjarnsholt T, Alhede MM, Alhede MM et al. The in vivo biofilm. Trends Microbiol 2013;21:466–74.

Blair JM, Richmond GE, Piddock LJ. Multidrug efflux pumps in Gram-negative bacteria and their role in antibiotic resistance. Futur Microbiol 2014;9:1165–77.

Blair JMA, Webber MA, Baylay AJ et al. Molecular mechanisms of antibiotic resistance. Nat Rev Microbiol 2015;13:42–51.

Blake KL, O’Neill AJ. Transposon library screening for identification of genetic loci participating in intrinsic susceptibility and acquired resistance to antistaphylococcal agents. J Antimicrob Chemother 2013;68:12–6.

Bolinger H, Kathariou S. The current state of macrolide resistance in Campylobacter spp.: Trends and impacts of resistance mechanisms. Appl Environ Microbiol 2017;83:e00416-17.

Bowen WH, Burne RA, Wu H et al. Oral biofilms: Pathogens, batrix, and polymicrobial interactions in microenvironments. Trends Microbiol 2018;26:229–42.
Brackman G, Coenye T. Quorum sensing inhibitors as anti-biofilm agents. *Curr Pharm Des* 2015;21:5–11.

Brauner A, Fridman O, Gefen O et al. Distinguishing between resistance, tolerance and persistence to antibiotic treatment. *Nat Rev Microbiol* 2016;14:320–30.

De Brucker K, Tan Y, Vints K et al. Fungal β-1,3-glucan increases ofloxacin tolerance of *Escherichia coli* in a polymicrobial *E. coli/Candida albicans* biofilm. *Antimicrob Agents Chemother* 2015;59:3052–8.

Cameron DR, Shan Y, Zalis EA et al. A genetic determinant of persister cell formation in bacterial pathogens. *J Bacteriol* 2018;200:e00303-18.

Cassini A, Högberg LD, Plachouras D et al. Attributable deaths and disability-adjusted life-years caused by infections with antibiotic-resistant bacteria in the EU and the European Economic Area in 2015: a population-level modelling analysis. *Lancet Infect Dis* 2018;19:56–66.

Castanheira M, Griffin MA, Deshpande LM et al. Detection of mcr-1 among *Escherichia coli* clinical isolates collected worldwide as part of the SENTRY antimicrobial surveillance program in 2014 and 2015. *Antimicrob Agents Chemother* 2016;60:5623–4.

CDC (2013). *Antibiotic resistance threats in the United States, 2013.* https://www.cdc.gov/drugresistance/threat-report-2013/pdf/ar-threats-2013-508.pdf (4 July 2019, date last accessed).

CDC (2017). *Sexually transmitted disease surveillance 2017: Gonorrhea.* https://www.cdc.gov/std/stats17/gonorrhea.htm (4 July 2019, date last accessed).

CDC (2018). *Shigellosis - Chapter 3 - 2018 Yellow Book | Travelers’ Health.* https://wwwnc.cdc.gov/travel/yellowbook/2018/infectious-diseases-related-to-travel/shigellosis (16 May 2019, date last accessed).

Chan K-G, Liu Y-C, Chang C-Y. Inhibiting N-acyl-homoserine lactone synthesis and quenching *Pseudomonas* quinolone quorum sensing to attenuate virulence. *Front Microbiol* 2015;6:1173.

Chiang W-C, Nilsson M, Jensen PØ et al. Extracellular DNA shields against aminoglycosides in *Pseudomonas aeruginosa* biofilms. *Antimicrob Agents Chemother* 2013;57:2352–61.

Conlon BP. *Staphylococcus aureus* chronic and relapsing infections: Evidence of a role for persister cells: An investigation of persister cells, their formation and their role in *S. aureus* disease. *BioEssays* 2014;36:991–6.

Conlon BP, Nakayasu ES, Fleck LE et al. Activated ClpP kills persisters and eradicates a chronic biofilm infection. *Nature* 2013;503:365–70.

Conlon BP, Rowe SE, Gandt AB et al. Persister formation in *Staphylococcus aureus* is associated with ATP depletion. *Nat Microbiol* 2016;1:16051.

Conlon BP, Rowe SE, Lewis K. Persister cells in biofilm associated infections. *Advances in Experimental Medicine and Biology.* 2015, 1–9.
Costa-Orlandi CB, Sardi JCO, Pitangui NS et al. Fungal biofilms and polymicrobial diseases. *J Fungi* 2017;3:E22.

Crumplin GC, Smith JT. Nalidixic acid and bacterial chromosome replication. *Nature* 1976;260:643–5.

D’Costa VM, King CE, Kalan L et al. Antibiotic resistance is ancient. *Nature* 2011;477:457–61.

Dalton T, Dowd SE, Wolcott RD et al. An in vivo polymicrobial biofilm wound infection model to study interspecies interactions. *PLoS One* 2011;6:e27317.

Desrousseaux C, Sautou V, Descamps S et al. Modification of the surfaces of medical devices to prevent microbial adhesion and biofilm formation. *J Hosp Infect* 2013;85:87–93.

Dolejska M, Villa L, Poirel L et al. Complete sequencing of an IncHI1 plasmid encoding the carbapenemase NDM-1, the ArmA 16S RNA methylase and a resistance-nodulation-cell division/multidrug efflux pump. *J Antimicrob Chemother* 2013;68:34–9.

Donné J, Dewilde S. The challenging world of biofilm physiology. *Adv Microb Physiol* 2015;67:235–92.

Doroshenko N, Tseng BS, Howlin RP et al. Extracellular DNA impedes the transport of vancomycin in *Staphylococcus epidermidis* biofilms preexposed to subinhibitory concentrations of vancomycin. *Antimicrob Agents Chemother* 2014;58:7273–82.

Dou Y, Song F, Guo F et al. *Acinetobacter baumannii* quorum-sensing signalling molecule induces the expression of drug-resistance genes. *Mol Med Rep* 2017;15:4061–8.

ECDC (2017a). *Factsheet about Invasive Haemophilus influenzae disease*. https://ecdc.europa.eu/en/invasive-haemophilus-influenzae-disease/facts (16 May 2019, date last accessed).

ECDC (2017b). *Factsheet about shigellosis*. https://ecdc.europa.eu/en/shigellosis/facts (16 May 2019, date last accessed).

EFSA. The European Union summary report on antimicrobial resistance in zoonotic and indicator bacteria from humans, animals and food in 2016. *EFSA J* 2018;16, DOI: 10.2903/j.efsa.2018.5182.

El-Halfawy OM, Valvano MA. Antimicrobial heteroresistance: an emerging field in need of clarity. *Clin Microbiol Rev* 2015;28:191–207.

Elias S, Banin E. Multi-species biofilms: living with friendly neighbors. *FEMS Microbiol Rev* 2012;36:990–1004.

Evans KC, Benomar S, Camuy-Vélez LA et al. Quorum-sensing control of antibiotic resistance stabilizes cooperation in *Chromobacterium violaceum*. *ISME J* 2018;12:1263–72.

Fernández L, Breidenstein EBM, Hancock REW. Creeping baselines and adaptive resistance to antibiotics. *Drug Resist Updat* 2011;14:1–21.
Fernandez L, Hancock REW. Adaptive and mutational resistance: Role of porins and efflux pumps in drug resistance. Clin Microbiol Rev 2012;25:661–81.

Filkins LM, O’Toole GA. Cystic fibrosis lung infections: Polymicrobial, complex, and hard to treat. PLoS Pathog 2015;11:e1005258.

Fisher RA, Gollan B, Helaine S. Persistent bacterial infections and persister cells. Nat Rev Microbiol 2017;15:453–64.

Flemming H-C, Wingender J. The biofilm matrix. Nat Rev Microbiol 2010;8:623–33.

Flemming H-C, Wingender J, Szewzyk U et al. Biofilms: an emergent form of bacterial life. Nat Rev Microbiol 2016;14:563–75.

Forsberg KJ, Patel S, Gibson MK et al. Bacterial phylogeny structures soil resistomes across habitats. Nature 2014;509:612–6.

Forslund K, Sunagawa S, Coelho LP et al. Metagenomic insights into the human gut resistome and the forces that shape it. BioEssays 2014;36:316–29.

Gabrilskia RA, Rumbaugh KP. Biofilm models of polymicrobial infection. Futur Microbiol 2015;10:1997–2015.

Gao W, Howden BP, Stinear TP. Evolution of virulence in Enterococcus faecium, a hospital-adapted opportunistic pathogen. Curr Opin Microbiol 2018;41:76–82.

Garcia LG, Lemaire S, Kahl BC et al. Antibiotic activity against small-colony variants of Staphylococcus aureus: review of in vitro, animal and clinical data. J Antimicrob Chemother 2013;68:1455–64.

Gardete S, Tomasz A. Mechanisms of vancomycin resistance in Staphylococcus aureus. J Clin Invest 2014;124:2836–40.

Gaynes R. The discovery of penicillin—New insights after more than 75 years of clinical use. Emerg Infect Dis 2017;23:849–53.

Giaouris E, Chorianopoulos N, Doulgeraki A et al. Co-Culture with Listeria monocytogenes within a dual-species biofilm community strongly increases resistance of Pseudomonas putida to benzalkonium chloride. PLoS One 2013;8:e77276.

Giaouris E, Heir E, Desvaux M et al. Intra- and inter-species interactions within biofilms of important foodborne bacterial pathogens. Front Microbiol 2015;6:841.

Gillings MR, Stokes HW. Are humans increasing bacterial evolvability? Trends Ecol Evol 2012;27:346–52.

Glinel K, Thebault P, Humblot V et al. Antibacterial surfaces developed from bio-inspired approaches. Acta Biomater 2012;8:1670–84.

Goderska K, Agudo Pena S, Alarcon T. Helicobacter pylori treatment: antibiotics or probiotics. Appl Microbiol Biotechnol 2018;102:1–7.
Golkar Z, Bagasra O, Pace DG. Bacteriophage therapy: a potential solution for the antibiotic resistance crisis. *J Infect Dev Ctries* 2014;8:129–36.

Gopu V, Meena CK, Shetty PH. Quercetin influences quorum sensing in food borne bacteria: *in-vitro* and *in-silico* evidence. *PLoS One* 2015;10:e0134684.

Grandclément C, Tannières M, Moréra S et al. Quorum quenching: role in nature and applied developments. *FEMS Microbiol Rev* 2016;40:86–116.

Grant SS, Hung DT. Persistent bacterial infections, antibiotic tolerance, and the oxidative stress response. *Virulence* 2013;4:273–83.

Grassi L, Di Luca M, Maisetta G et al. Generation of persister cells of *Pseudomonas aeruginosa* and *Staphylococcus aureus* by chemical treatment and evaluation of their susceptibility to membrane-targeting agents. *Front Microbiol* 2017;8:1917.

Hall CW, Mah T-F. Molecular mechanisms of biofilm-based antibiotic resistance and tolerance in pathogenic bacteria. *FEMS Microbiol Rev* 2017;41:276–301.

Harding CM, Hennon SW, Feldman MF. Uncovering the mechanisms of *Acinetobacter baumannii* virulence. *Nat Rev Microbiol* 2017;16:91–102.

Hasan J, Crawford RJ, Ivanova EP. Antibacterial surfaces: The quest for a new generation of biomaterials. *Trends Biotechnol* 2013;31:295–304.

Hassoun A, Linden PK, Friedman B. Incidence, prevalence, and management of MRSA bacteremia across patient populations—a review of recent developments in MRSA management and treatment. *Crit Care* 2017;21:211.

Hawver LA, Jung SA, Ng W-L. Specificity and complexity in bacterial quorum-sensing systems. *FEMS Microbiol Rev* 2016;40:738–52.

He J, Chen J, Hu G et al. Immobilization of an antimicrobial peptide on silicon surface with stable activity by click chemistry. *J Mater Chem B* 2018;6:68–74.

Hirakawa H, Tomita H. Interference of bacterial cell-to-cell communication: a new concept of antimicrobial chemotherapy breaks antibiotic resistance. *Front Microbiol* 2013;4:114.

Høiby N, Bjarnsholt T, Moser C et al. ESCMID guideline for the diagnosis and treatment of biofilm infections 2014. *Clin Microbiol Infect* 2015;21:S1–25.

Horne D, Tomasz A. Tolerant response of *Streptococcus sanguis* to beta-lactams and other cell wall inhibitors. *Antimicrob Agents Chemother* 1977;11:888–96.

Hossain MA, Lee SJ, Park NH et al. Impact of phenolic compounds in the acyl homoserine lactone-mediated quorum sensing regulatory pathways. *Sci Rep* 2017;7:10618.

Hsu LC, Fang J, Borca-Tasciuc DA et al. Effect of micro- and nanoscale topography on the adhesion of bacterial cells to solid surfaces. *Appl Environ Microbiol* 2013;79:2703–12.
Huang K, Lee BP, Ingram DR et al. Synthesis and characterization of self-assembling block copolymers containing bioadhesive end groups. Biomacromolecules 2002;3:397–406.

Johnson L, Horsman SR, Charron-Mazenod L et al. Extracellular DNA-induced antimicrobial peptide resistance in Salmonella enterica serovar Typhimurium. BMC Microbiol 2013;13:115.

Jorge P, Alves D, Pereira MO. Catalysing the way towards antimicrobial effectiveness: a systematic analysis and a new online resource for antimicrobial enzyme combinations against Pseudomonas aeruginosa and Staphylococcus aureus. Int J Antimicrob Agents 2019;53:598–605.

Kalia VC, Patel SKS, Kang YC et al. Quorum sensing inhibitors as antipathogens: biotechnological applications. Biotechnol Adv 2019;37:68–90.

Kang T, Banquy X, Heo J et al. Mussel-inspired anchoring of polymer loops that provide superior surface lubrication and antifouling properties. ACS Nano 2016;10:930–7.

Kaplan JB, Ragunath C, Ramasubbu N et al. Detachment of Actinobacillus actinomycetemcomitans biofilm cells by an endogenous β-hexosaminidase activity. J Bacteriol 2003;185:4693–8.

Khameneh B, Diab R, Ghazvini K et al. Breakthroughs in bacterial resistance mechanisms and the potential ways to combat them. Microb Pathog 2016;95:32–42.

Khandekar S, Liebens V, Fauvart M et al. The putative De-N-acetylase DnpA contributes to intracellular and biofilm-associated persistence of Pseudomonas aeruginosa exposed to fluoroquinolones. Front Microbiol 2018;9:1455.

Kim C, Kim J, Park HY et al. Furanone derivatives as quorum-sensing antagonists of Pseudomonas aeruginosa. Appl Microbiol Biotechnol 2008;80:37–47.

Kim S-Y, Lee S-K, Park M-S et al. Analysis of the fluoroquinolone antibiotic resistance mechanism of Salmonella enterica isolates. J Microbiol Biotechnol 2016;26:1605–12.

Knecht L, O’Connor G, Mittal R et al. Serotonin activates bacterial quorum sensing and enhances the virulence of Pseudomonas aeruginosa in the host. EBioMedicine 2016;9:161–169.

Korgaonkar A, Trivedi U, Rumbaugh KP et al. Community surveillance enhances Pseudomonas aeruginosa virulence during polymicrobial infection. Proc Natl Acad Sci U S A 2013;110:1059–64.

Kostakioti M, Hadjifrangiskou M, Hultgren SJ. Bacterial biofilms: Development, dispersal, and therapeutic strategies in the dawn of the postantibiotic era. Cold Spring Harb Perspect Med 2013;3:a010306.

Kumar A, Alam A, Rani M et al. Biofilms: Survival and defense strategy for pathogens. Int J Med Microbiol 2017;307:481–9.

Kumar A, Schweizer H. Bacterial resistance to antibiotics: Active efflux and reduced uptake. Adv Drug Deliv Rev 2005;57:1486–513.
Kumar P, Kizhakkedathu JN, Straus SK. Antimicrobial peptides: Diversity, mechanism of action and strategies to improve the activity and biocompatibility in vivo. Biomolecules 2018;8:4.

Kumar Shukla S, Rao TS. Dispersal of Bap-mediated Staphylococcus aureus biofilm by proteinase K. J Antibiot 2013;66:55–60.

de la Fuente-Núñez C, Reffuveille F, Fernández L et al. Bacterial biofilm development as a multicellular adaptation: antibiotic resistance and new therapeutic strategies. Curr Opin Microbiol 2013;16:580–9.

Lade H, Paul D, Kweon JH. N-acyl homoserine lactone-mediated quorum sensing with special reference to use of quorum quenching bacteria in membrane biofouling control. BioMed Res Int 2014;2014:162584.

Landecker H. Antibiotic resistance and the biology of history. Body Soc 2016;22:19–52.

Langton KP, Henderson PJF, Herbert RB. Antibiotic resistance: multidrug efflux proteins, a common transport mechanism? Nat Prod Rep 2005;22:439.

LaSarre B, Federle MJ. Exploiting quorum sensing to confuse bacterial pathogens. Microbiol Mol Biol Rev 2013;77:73–111.

Lavigne J-P, Sotto A, Nicolas-Chanoine M-H et al. An adaptive response of Enterobacter aerogenes to imipenem: regulation of porin balance in clinical isolates. Int J Antimicrob Agents 2013;41:130–6.

Laxminarayan R, Heymann DL. Challenges of drug resistance in the developing world. BMJ 2012;344:e1567–e1567.

Lebeaux D, Ghigo J-M, Beloin C. Biofilm-related infections: bridging the gap between clinical management and fundamental aspects of recalcitrance toward antibiotics. Microbiol Mol Biol Rev 2014;78:510–43.

Lee C-R, Lee JH, Park M et al. Biology of Acinetobacter baumannii: Pathogenesis, antibiotic resistance mechanisms, and prospective treatment options. Front Cell Infect Microbiol 2017;7:55.

Lee KWK, Periasamy S, Mukherjee M et al. Biofilm development and enhanced stress resistance of a model, mixed-species community biofilm. ISME J 2014;8:894–907.

Li XZ, Livermore DM, Nikaido H. Role of efflux pump(s) in intrinsic resistance of Pseudomonas aeruginosa: resistance to tetracycline, chloramphenicol, and norfloxacin. Antimicrob Agents Chemother 1994;38:1732–41.

Li XZ, Ma D, Livermore DM et al. Role of efflux pump(s) in intrinsic resistance of Pseudomonas aeruginosa: active efflux as a contributing factor to beta-lactam resistance. Antimicrob Agents Chemother 1994;38:1742–52.

Li XZ, Nikaido H, Poole K. Role of mexA-mexB-oprM in antibiotic efflux in Pseudomonas aeruginosa. Antimicrob Agents Chemother 1995;39:1948–53.
Li Y-H, Tian X. Quorum sensing and bacterial social interactions in biofilms. *Sensors (Basel)* 2012;12:2519–38.

Liu A, Tran L, Becket E *et al.* Antibiotic sensitivity profiles determined with an *Escherichia coli* gene knockout collection: generating an antibiotic bar code. *Antimicrob Agents Chemother* 2010;54:1393–403.

Lopes SP, Ceri H, Azevedo NF *et al.* Antibiotic resistance of mixed biofilms in cystic fibrosis: impact of emerging microorganisms on treatment of infection. *Int J Antimicrob Agents* 2012;40:260–3.

Lushniak BD. Antibiotic resistance: a public health crisis. *Public Heal Rep* 2014;129:314–6.

Luyt C-E, Bréchot N, Trouillet J-L *et al.* Antibiotic stewardship in the intensive care unit. *Crit Care* 2014;18:480.

Magalhães AP, Lopes SP, Pereira MO. Insights into cystic fibrosis polymicrobial consortia: the role of species interactions in biofilm development, phenotype, and response to in-use antibiotics. *Front Microbiol* 2017;7:1–11.

Magana M, Sereti C, Ioannidis A *et al.* Options and limitations in clinical investigation of bacterial biofilms. *Clin Microbiol Rev* 2018;31:e00084-16.

Marques CNH, Davies DG, Sauer K. Control of biofilms with the fatty acid signaling molecule cis-2-decenolic acid. *Pharmaceuticals* 2015;8:816–35.

Marshall BM, Levy SB. Food animals and antimicrobials: Impacts on human health. *Clin Microbiol Rev* 2011;24:718–33.

Martens E, Demain AL. The antibiotic resistance crisis, with a focus on the United States. *J Antibi* 2017;70:520–6.

Mashburn LM, Jett AM, Akins DR *et al.* *Staphylococcus aureus* serves as an iron source for *Pseudomonas aeruginosa* during in vivo coculture. *J Bacteriol* 2005;187:554–66.

McCann CM, Christgen B, Roberts JA *et al.* Understanding drivers of antibiotic resistance genes in High Arctic soil ecosystems. *Environ Int* 2019;125:497–504.

El Meouche I, Dunlop MJ. Heterogeneity in efflux pump expression predisposes antibiotic-resistant cells to mutation. *Science* 2018;362:686–90.

Michael CA, Dominey-Howes D, Labbate M. The antimicrobial resistance crisis: causes, consequences, and management. *Front Public Heal* 2014;2:145.

Millezi FM, Pereira MO, Batista NN *et al.* Susceptibility of monospecies and dual-species biofilms of *Staphylococcus aureus* and *Escherichia coli* to essential oils. *J Food Saf* 2012;32:351–9.

Mitchell G, Séguin DL, Asselin A-E *et al.* *Staphylococcus aureus* sigma B-dependent emergence of small-colony variants and biofilm production following exposure to *Pseudomonas aeruginosa* 4-hydroxy-2-heptylquinoline-N-oxide. *BMC Microbiol* 2010;10:33.
Miyaue S, Suzuki E, Komiyama Y et al. Bacterial memory of persisters: Bacterial persister cells can retain their phenotype for days or weeks after withdrawal from colony–biofilm culture. Front Microbiol 2018;9:1396.

Monaco M, Pimentel de Araujo F, Cruciani M et al. Worldwide epidemiology and antibiotic resistance of Staphylococcus aureus. Curr Top Microbiol Immunol 2016;409:21–56.

Morgan DJ, Okeke IN, Laxminarayan R et al. Non-prescription antimicrobial use worldwide: a systematic review. Lancet Infect Dis 2011;11:692–701.

Motta SS, Cluzel P, Aldana M. Adaptive resistance in bacteria requires epigenetic inheritance, genetic noise, and cost of efflux pumps. PLoS One 2015;10:e0118464.

Munita JM, Arias CA. Mechanisms of antibiotic resistance. Microbiol Spectr 2016;4:481–511.

Murray JL, Connell JL, Stacy A et al. Mechanisms of synergy in polymicrobial infections. J Microbiol 2014;52:188–99.

Nature Editorial. The antibiotic alarm. Nature 2013;495:141.

Nikaido H. Molecular basis of bacterial outer membrane permeability revisited. Microbiol Mol Biol Rev 2003;67:593–656.

Nikaido H. Multidrug resistance in bacteria. Annu Rev Biochem 2009;78:119–46.

O’Neill J (2014). Antimicrobial Resistance: Tackling a crisis for the health and wealth of nations. The Review on Antimicrobial Resistance. https://amr-review.org/sites/default/files/AMR20Review20Paper20-%20Tackling%20a%20crisis%20for%20the%20health%20and%20wealth%20of%20nations_1.pdf (14 January 2019, date last accessed).

O’Toole GA, Ha D-G. c-di-GMP and its effects on biofilm formation and dispersion: a Pseudomonas aeruginosa review. Microbiol Spectr 2015;3:MB-0003-2014.

Okshevsky M, Meyer RL. The role of extracellular DNA in the establishment, maintenance and perpetuation of bacterial biofilms. Crit Rev Microbiol 2015;41:341–52.

Olsen I. Biofilm-specific antibiotic tolerance and resistance. Eur J Clin Microbiol Infect Dis 2015;34:877–86.

Orazi G, O’Toole GA. Pseudomonas aeruginosa alters Staphylococcus aureus sensitivity to vancomycin in a biofilm model of cystic fibrosis infection. MBio 2017;8:e00873-17.

Pallecchi L, Bartoloni A, Paradisi F et al. Antibiotic resistance in the absence of antimicrobial use: mechanisms and implications. Expert Rev Anti Infect Ther 2008;6:725–32.

Pang Z, Raudonis R, Glick BR et al. Antibiotic resistance in Pseudomonas aeruginosa: mechanisms and alternative therapeutic strategies. Biotechnol Adv 2019;37:177–92.

Parrino B, Schillaci D, Carnevale I et al. Synthetic small molecules as anti-biofilm agents in the
struggle against antibiotic resistance. *Eur J Med Chem* 2019;**161**:154–78.

Passos da Silva D, Schofield M, Parsek M *et al.* An update on the sociomicrobiology of quorum sensing in Gram-negative biofilm development. *Pathogens* 2017;**6**:51.

Pastar I, Nusbaum AG, Gil J *et al.* Interactions of methicillin resistant *Staphylococcus aureus* USA300 and *Pseudomonas aeruginosa* in polymicrobial wound infection. *PLoS One* 2013;**8**:e56846.

Pawlowski AC, Wang W, Koteva K *et al.* A diverse intrinsic antibiotic resistome from a cave bacterium. *Nat Commun* 2016;**7**:13803.

Pechère J-C, Köhler T. Patterns and modes of beta-lactam resistance in *Pseudomonas aeruginosa*. *Clin Microbiol Infect* 1999;**5 Suppl 1**:S15–8.

Pérez-Pérez M, Jorge P, Pérez Rodríguez G *et al.* Quorum sensing inhibition in *Pseudomonas aeruginosa* biofilms: new insights through network mining. *Biofouling* 2017;**33**:128–42.

Perez AC, Pang B, King LB *et al.* Residence of *Streptococcus pneumoniae* and *Moraxella catarrhalis* within polymicrobial biofilm promotes antibiotic resistance and bacterial persistence in vivo. *Pathog Dis* 2014;**70**:280–8.

Perron GG, Whyte L, Turnbaugh PJ *et al.* Functional characterization of bacteria isolated from ancient arctic soil exposes diverse resistance mechanisms to modern antibiotics. *PLoS One* 2015;**10**:e0069533.

Perry JA, Westman EL, Wright GD. The antibiotic resistome: what’s new? *Curr Opin Microbiol* 2014;**21**:45–50.

Petchiappan A, Chatterji D. Antibiotic resistance: Current perspectives. *ACS Omega* 2017;**2**:7400–9.

Peters BM, Jabra-Rizk MA, O’May GA *et al.* Polymicrobial interactions: Impact on pathogenesis and human disease. *Clin Microbiol Rev* 2012;**25**:193–213.

Peterson E, Kaur P. Antibiotic resistance mechanisms in bacteria: Relationships between resistance determinants of antibiotic producers, environmental bacteria, and clinical pathogens. *Front Microbiol* 2018;**9**:2928.

Pires D, Melo L, Vilas Boas D *et al.* Phage therapy as an alternative or complementary strategy to prevent and control biofilm-related infections. *Curr Opin Microbiol* 2017;**39**:48–56.

Pompilio A, Crocetta V, De Nicola S *et al.* Cooperative pathogenicity in cystic fibrosis: *Stenotrophomonas maltophilia* modulates *Pseudomonas aeruginosa* virulence in mixed biofilm. *Front Microbiol* 2015;**6**:951.

Poole K. Efflux-mediated antimicrobial resistance. *J Antimicrob Chemother* 2005;**56**:20–51.

Potron A, Poirel L, Nordmann P. Emerging broad-spectrum resistance in *Pseudomonas aeruginosa* and *Acinetobacter baumannii*: Mechanisms and epidemiology. *Int J Antimicrob Agents* 2015;**45**:568–85.
Proctor RA, Kriegeskorte A, Kahl BC et al. *Staphylococcus aureus* Small Colony Variants (SCVs): a road map for the metabolic pathways involved in persistent infections. *Front Cell Infect Microbiol* 2014;4:99.

Ragland SA, Criss AK. From bacterial killing to immune modulation: Recent insights into the functions of lysozyme. *PLoS Pathog* 2017;13:e1006512.

Ramsey DM, Wozniak DJ. Understanding the control of *Pseudomonas aeruginosa* alginate synthesis and the prospects for management of chronic infections in cystic fibrosis. *Mol Microbiol* 2005;56:309–22.

Ramsey MM, Rumbaugh KP, Whiteley M. Metabolite cross-feeding enhances virulence in a model polymicrobial infection. *PLoS Pathog* 2011;7:e1002012.

Randall CP, Mariner KR, Chopra I et al. The target of daptomycin is absent from *Escherichia coli* and other Gram-negative pathogens. *Antimicrob Agents Chemother* 2013;57:637–9.

Rawat D, Nair D. Extended-spectrum β-lactamases in Gram negative bacteria. *J Glob Infect Dis* 2010;2:263–74.

Read AF, Woods RJ. Antibiotic resistance management. *Evol Med Public Heal* 2014;2014:147–147.

Rémy B, Mion S, Plener L et al. Interference in bacterial quorum sensing: a biopharmaceutical perspective. *Front Pharmacol* 2018;9:203.

Rice LB. Mechanisms of resistance and clinical relevance of resistance to β-lactams, glycopeptides, and fluoroquinolones. *Mayo Clin Proc* 2012;87:198–208.

Rice PA, Shafer WM, Ram S et al. *Neisseria gonorrhoeae*: Drug resistance, mouse models, and vaccine development. *Annu Rev Microbiol* 2017;71:665–86.

Richter MF, Drown BS, Riley AP et al. Predictive compound accumulation rules yield a broad-spectrum antibiotic. *Nature* 2017;545:299–304.

Roca I, Akova M, Baquero F et al. The global threat of antimicrobial resistance: Science for intervention. *New Microbes New Infect* 2015;6:22–9.

Rossolini GM, Arena F, Pecile P et al. Update on the antibiotic resistance crisis. *Curr Opin Pharmacol* 2014;18:56–60.

Ryan RP, Fouhy Y, Garcia BF et al. Interspecies signalling via the *Stenotrophomonas maltophilia* diffusible signal factor influences biofilm formation and polymyxin tolerance in *Pseudomonas aeruginosa*. *Mol Microbiol* 2008;68:75–86.

Sahukhal GS, Pandey S, Elasri MO. msaABCR operon is involved in persister cell formation in *Staphylococcus aureus*. *BMC Microbiol* 2017;17:218.

Sambanthamoorthy K, Luo C, Pattabiraman N et al. Identification of small molecules inhibiting diguanylate cyclases to control bacterial biofilm development. *Biofouling* 2014;30:17–28.
Sanchez-Romero MA, Casadesus J. Contribution of phenotypic heterogeneity to adaptive antibiotic resistance. *Proc Natl Acad Sci U S A* 2014;111:355–60.

Sanchez-Vizuete P, Le Coq D, Bridier A et al. Identification of ypqP as a new *Bacillus subtilis* biofilm determinant that mediates the protection of *Staphylococcus aureus* against antimicrobial agents in mixed-species communities. *Appl Environ Microbiol* 2015;81:109–18.

Sandoval-Motta S, Aldana M. Adaptive resistance to antibiotics in bacteria: a systems biology perspective. *Wiley Interdiscip Rev Syst Biol Med* 2016;8:253–67.

Schaible B, Taylor CT, Schaffer K. Hypoxia increases antibiotic resistance in *Pseudomonas aeruginosa* through altering the composition of multidrug efflux pumps. *Antimicrob Agents Chemother* 2012;56:2114–8.

Schertzer JW, Boulette ML, Whiteley M. More than a signal: Non-signaling properties of quorum sensing molecules. *Trends Microbiol* 2009;17:189–95.

Schroeder M, Brooks BD, Brooks AE. The complex relationship between virulence and antibiotic resistance. *Genes (Basel)* 2017;8:E39.

Schwering M, Song J, Louie M et al. Multi-species biofilms defined from drinking water microorganisms provide increased protection against chlorine disinfection. *Biofouling* 2013;29:917–28.

Shan Y, Brown Gandt A, Rowe SE et al. ATP-dependent persister formation in *Escherichia coli*. *MBio* 2017;8:e02267-16.

Smith PA, Koehler MFT, Girgis HS et al. Optimized arylomycins are a new class of Gram-negative antibiotics. *Nature* 2018;561:189–94.

Smith T, Wolff KA, Nguyen L. Molecular biology of drug resistance in *Mycobacterium tuberculosis*. *Curr Top Microbiol Immunol* 2012;374:53–80.

Sriram KB, Cox AJ, Clancy RL et al. Nontypeable *Haemophilus influenzae* and chronic obstructive pulmonary disease: a review for clinicians. *Crit Rev Microbiol* 2018;44:125–42.

Stacy A, McNally L, Darch SE et al. The biogeography of polymicrobial infection. *Nat Rev Microbiol* 2016;14:93–105.

Stewart PS. Biophysics of biofilm infection. *Pathog Dis* 2014;70:212–8.

Stokes HW, Gillings MR. Gene flow, mobile genetic elements and the recruitment of antibiotic resistance genes into Gram-negative pathogens. *FEMS Microbiol Rev* 2011;35:790–819.

Sugimoto S, Sato F, Miyakawa R et al. Broad impact of extracellular DNA on biofilm formation by clinically isolated methicillin-resistant and -sensitive strains of *Staphylococcus aureus*. *Sci Rep* 2018;8:2254.

Tangden T, Adler M, Cars O et al. Frequent emergence of porin-deficient subpopulations with reduced carbapenem susceptibility in ESBL-producing *Escherichia coli* during exposure to
ertapenem in an in vitro pharmacokinetic model. J Antimicrob Chemother 2013;68:1319–26.

Tata M, Wolfinger MT, Amman F et al. RNASeq based transcriptional profiling of Pseudomonas aeruginosa PA14 after short- and long-term anoxic cultivation in synthetic cystic fibrosis sputum medium. PLoS One 2016;11:e0147811.

Taylor PK, Yeung ATY, Hancock REW. Antibiotic resistance in Pseudomonas aeruginosa biofilms: towards the development of novel anti-biofilm therapies. J Biotechnol 2014;191:121–30.

Turkina M V, Vikström E. Bacteria-host crosstalk: Sensing of the quorum in the context of Pseudomonas aeruginosa infections. J Innate Immun 2018;14:1–17.

Valentini M, Filloux A. Biofilms and Cyclic di-GMP (c-di-GMP) signaling: Lessons from Pseudomonas aeruginosa and other bacteria. J Biol Chem 2016;291:12547–55.

Vega LM, Mathieu J, Yang Y et al. Nickel and cadmium ions inhibit quorum sensing and biofilm formation without affecting viability in Burkholderia multivorans. Int Biodeter Biodegr 2014;91:82–7.

Ventola CL. The antibiotic resistance crisis: Part 1: Causes and threats. P T 2015;40:277–83.

Vulin C, Leimer N, Huemer M et al. Prolonged bacterial lag time results in small colony variants that represent a sub-population of persisters. Nat Commun 2018;9:4074.

Wang C, Sauvageau D, Elias A. Immobilization of active bacteriophages on polyhydroxyalkanoate surfaces. ACS Appl Mater Interfaces 2016;8:1128–38.

Wang LH, Weng LX, Dong YH et al. Specificity and enzyme kinetics of the quorum-quenching N-acyl homoserine lactone lactonase (AHL-lactonase). J Biol Chem 2004;279:13645–51.

Watkins RR, David MZ, Salata RA. Current concepts on the virulence mechanisms of meticillin-resistant Staphylococcus aureus. J Med Microbiol 2012;61:1179–93.

Whitchurch CB, Tolker-Nielsen T, Ragas PC et al. Extracellular DNA required for bacterial biofilm formation. Science (80- ) 2002;295:1487.

WHO (2014). Immunization, vaccines and biologicals: Pneumococcal disease. https://www.who.int/immunization/diseases/pneumococcal/ (4 July 2019, date last accessed).

WHO (2017a). Global priority list of antibiotic-resistant bacteria to guide research, discovery, and development of new antibiotics. https://www.who.int/medicines/publications/WHO-PPL-Short_Summary_25Feb-ET_NM_WHO.pdf (13 May 2019, date last accessed).

WHO (2017b). Guidelines for the prevention and control of carbapenem-resistant Enterobacteriaceae, Acinetobacter baumannii and Pseudomonas aeruginosa in health care facilities. https://www.ncbi.nlm.nih.gov/books/NBK493062/ (4 July 2019, date last accessed).

WHO (2018a). Global tuberculosis report 2018. https://www.who.int/tb/publications/global_report/en/ (4 July 2019, date last accessed).
WHO (2018b). *Salmonella (non-typhoidal).* [https://www.who.int/news-room/fact-sheets/detail/salmonella-(non-typhoidal)](https://www.who.int/news-room/fact-sheets/detail/salmonella-(non-typhoidal)) (15 May 2019, date last accessed).

Wilton M, Charron-Mazenod L, Moore R *et al.* Extracellular DNA acidifies biofilms and induces aminoglycoside resistance in *Pseudomonas aeruginosa.* *Antimicrob Agents Chemother* 2016;60:544–53.

Wolcott R, Costerton JW, Raoult D *et al.* The polymicrobial nature of biofilm infection. *Clin Microbiol Infect* 2013;19:107–12.

Wozniak A, Villagra NA, Undabarrena A *et al.* Porin alterations present in non-carbapenemase-producing Enterobacteriaceae with high and intermediate levels of carbapenem resistance in Chile. *J Med Microbiol* 2012;61:1270–9.

Xu W, DeJesus MA, Rücker N *et al.* Chemical genetic interaction profiling reveals determinants of intrinsic antibiotic resistance in *Mycobacterium tuberculosis.* *Antimicrob Agents Chemother* 2017;61:e01334-17.

Yadav MK, Park SW, Chae SW *et al.* Sinefungin, a natural nucleoside analogue of S-adenosylmethionine, inhibits *Streptococcus pneumoniae* biofilm growth. *BioMed Res Int* 2014;2014:156987.

Zaltsman N, Ionescu AC, Weiss EI *et al.* Surface-modified nanoparticles as anti-biofilm filler for dental polymers. *PLoS One* 2017;12:e0189397.

Zhu L, Lin J, Ma J *et al.* Triclosan resistance of *Pseudomonas aeruginosa* PAO1 is due to FabV, a triclosan-resistant enoyl-acyl carrier protein reductase. *Antimicrob Agents Chemother* 2010;54:689–98.
Figure 1. Biofilm resistance and tolerance mechanisms. Main resistance and tolerance mechanisms (in black) are characterized as intrinsic, acquired, or adaptive. Different geometric forms and colors denote different bacterial species. Nutrient and oxygen gradients are illustrated as a downwards triangle going from high (green top) to low (red bottom) concentration. Inability of antimicrobials to act upon the cell if represented as a red cross sign. Dormant or persister cell is depicted in gray. Legend: SCV – small-colony variant.
Table 1. The top MDR bacteria: WHO categorization and key features.

| WHO categorization | Bacterial pathogens | Key features | References |
|--------------------|---------------------|--------------|------------|
| Critical           | *Acinetobacter baumannii*, carbapenem-resistant | Most associated with HAIs worldwide, accounting for up to 20% of ICU infections worldwide. Causes pneumonia and bloodstream and wound infections, particularly in mechanically ventilated patients. Around 45% isolates are MDR, including resistance to last-resort carbapenems most often linked to the production of carbapenemases. | Potron, Poirel and Nordmann 2015; Harding, Hennon and Feldman 2017; Lee *et al.* 2017 |
|                    | *Pseudomonas aeruginosa*, carbapenem-resistant | Common cause of HAIs, including pneumonia, bloodstream, urinary tract, and surgical-site infections. Carbapenem resistance mostly related to porin (OprD) deficiency. Invasive isolates resistant to carbapenems were 17.8% in Europe (2015) and 19.2% in the USA (2014). | Potron, Poirel and Nordmann 2015; WHO 2017b |
| Enterobacteriaceae, carbapenem-resistant, 3rd generation cephalosporin-resistant | Enterobacteriaceae include *K. pneumonia*, *E. coli*, *Enterobacter* spp., *Serratia* spp., *Proteus* spp., *Providencia* spp, and *Morganella* spp. *K. pneumoniae* invasive isolates resistant to carbapenems were reported from all WHO regions, with some countries reporting up to 50%. Human isolates resistant to colistin, a last resort antimicrobial against carbapenem-resistant Enterobacteriaceae, were already reported. 10–20 % of Enterobacteriaceae isolated in the USA are resistant to ceftazidime. | Arizpe *et al.* 2016; Castanheira *et al.* 2016; WHO 2017b |
|                    | *Mycobacteria tuberculosis* | *M. tuberculosis* infection is the precursor to tuberculosis disease, responsible for 1.5 million deaths/year. Aerial dissemination, with infection typically occurring in the lungs. Sometimes treatable with first-line drugs (isoniazid, rifampicin) but mostly resistant to a several antibiotics (fluoroquinolones), and to second-line injectable drugs (amikacin, capreomycin, and kanamycin). | WHO 2018 |
| **Organism** | **Characteristics** | **References** |
|--------------|---------------------|---------------|
| *Enterococcus faecium*, vancomycin-resistant | Most commonly isolated Gram-positive nosocomial pathogen worldwide with highly flexible genome that enables rapid adaption. Vancomycin-resistant isolates rose from 0% to more than 80% from 1980 to 2007, in the USA. Vancomycin resistance arises from reduced vancomycin-binding affinity, involving alterations in the peptidoglycan synthesis pathway. | Arias and Murray 2012; Gao, Howden and Stinear 2018 |
| *Staphylococcus aureus*, methicillin-resistant (MRSA) | Among the most frequent of all antibiotic-resistant threats and leading cause of bacteremia. Outstanding versatility in adapting to different epidemiological settings (healthcare, community, animal). Characteristically MDR, with infections spreading across the globe. Infections commonly involve the skin, soft tissue, bone, joints, and indwelling catheters or prosthetic devices. | Monaco et al. 2016; Hassoun, Linden and Friedman 2017 |
| *Helicobacter pylori*, clarithromycin-resistant | Most successful human gastric pathogen able to resist stomach acids, colonizing over 50% of the population. Related to gastritis, peptic ulcers, gastric adenocarcinoma, iron deficiency anemia, idiopathic thrombocytopenic purpura, and vitamin B12 deficiency. Sequential, bismuth quadruple, and non-bismuth quadruple therapies seem effective in high clarithromycin-resistance countries. | Alba, Blanco and Alarcón 2017; Goderska, Agudo Pena and Alarcon 2018 |
| *Campylobacter spp.*, fluoroquinolone-resistant | Leading cause of foodborne illnesses, majorly gastroenteritis, primarily caused by *Campylobacter jejuni*. Antibiotic treatment is only recommended in vulnerable patients, as the young, the | Bolinger and Kathariou 2017 |
| Pathogen | Description | Antibiotic Treatment | References |
|----------|-------------|----------------------|------------|
| **Salmonella spp., fluoroquinolone-resistant** | Leading cause of foodborne illnesses/diarrheal diseases, namely gastroenteritis. Antibiotic treatment is only recommended in vulnerable patients, as the young, the elderly, and patients with weakened immunity. | Kim et al. 2016; WHO 2018b |
| **Neisseria gonorrhoeae, 3rd generation cephalosporin-resistant, fluoroquinolone-resistant** | Causes gonorrhea, an obligate human infection, usually transmitted during sexual activity, often resulting in urethritis in men and cervicitis in women. Gonorrhea is rising, with 18.6% increase during 2016–2017 and 75.2% increase since 2009 in the USA. Asymptomatic men (two-thirds of infected men) constituting the principal source of dissemination. | CDC 2017; Rice et al. 2017 |
| **Streptococcus pneumoniae, penicillin-non-susceptible** | Encapsulated bacteria causes meningitis, septicaemia, and pneumonia, but also milder infections, such as sinusitis and otitis media. Major cause of morbidity and mortality worldwide, mainly in poor countries and in children under the age of two. There are two available vaccines that target the most prevalent serotypes. | WHO 2014 |
| **Medium** | Serotype b, an obligate human pathogen, is the most pathogenic, responsible for respiratory infections, ocular infection, sepsis, and meningitis. Leading worldwide cause of meningitis morbidity and mortality in unimmunised populations. Highly related to Chronic Obstructive Pulmonary Disease, a leading cause of morbidity and mortality worldwide. 3rd generation cephalosporins are the empiric treatment of choice. | ECDC 2017; Sriram et al. 2018 |
| **Shigella spp., fluoroquinolone-resistant** | Causes Shigellosis, a major cause of diarrhoea affecting mainly children under the age of five. | (ECDC 2017b; CDC 2018) |
| | | Between 80–165 million cases of shigellosis occur annually worldwide, majorly in developing countries. | |

Note: HAIs – hospital acquired infections; ICU – intensive care unit.
Table 2. Examples of inter-species interactions leading to increased AMR in polymicrobial biofilms for selected antimicrobial agents.

| Antimicrobial   | Species                                                      | Interaction outcome                                                                 | References         |
|-----------------|--------------------------------------------------------------|-------------------------------------------------------------------------------------|--------------------|
| Amoxicillin     | *Moraxella catarrhalis, Streptococcus pneumoniae*            | *M. catarrhalis* secreted β-lactamases protected *S. pneumonia* from amoxicillin treatment. | Perez et al., 2014 |
| Ampicillin      | *Haemophilus influenzae*, *Moraxella catarrhalis*            | *M. catarrhalis* secreted β-lactamases protected *H. influenzae* from ampicillin treatment. | Armbruster et al. 2010 |
| Azithromycin    | *Moraxella catarrhalis, Streptococcus pneumoniae*            | *S. pneumonia* protected *M. catarrhalis* from azithromycin treatment a signaling molecule AI-2 independent mechanism. | Perez et al., 2014 |
| Benzalkonium chloride | *Listeria monocytogenes*, *Pseudomonas putida*            | *L. monocytogenes* increased *P. putida* resistance to benzalkonium chloride.    | Giaouris et al., 2013 |
| Cefotaxime      | *Dolosigranulum pigrum*, *Inquilinus limosus*, *Pseudomonas aeruginosa* | *P. aeruginosa* increased *I. limosus* and *D. pigrum* resistance to cefotaxime. | Lopes et al., 2012  |
| Chloramphenicol | *Dolosigranulum pigrum*, *Inquilinus limosus*, *Pseudomonas aeruginosa* | *P. aeruginosa* increased *I. limosus* and *D. pigrum* resistance to chloramphenicol. | Lopes et al., 2012  |
| Chlorine        | *Enterobacteriaceae cloacae*, *Escherichia coli*, *Pseudomonas aeruginosa*, *Stenotrophomonas maltophilia* | Multi-species biofilms of the four bacteria displayed increased resistance to chlorine. | Schwering et al. 2013 |
| Ciprofloxacin   | *Inquilinus limosus*, *Pseudomonas aeruginosa*, *Staphylococcus aureus*, *Stenotrophomonas maltophilia* | *P. aeruginosa* increased *S. aureus*, *I. limosus*, and *S. maltophilia* resistance to ciprofloxacin. | Magalhães et al., 2017 |
| Clarithromycin  | *Haemophilus influenzae*, *Moraxella catarrhalis*            | *H. influenzae* signalling molecule AI-2 induced *M. catarrhalis* resistance to clarithromycin. | Armbruster et al. 2010 |
| Clindamycin     | *Dolosigranulum pigrum*, *Inquilinus limosus*, *Pseudomonas aeruginosa* | *P. aeruginosa* increased *I. limosus* and *D. pigrum* resistance to clindamycin. | Lopes et al., 2012  |
| Colistin        | *Pseudomonas aeruginosa*, *Stenotrophomonas maltophilia*    | *S. maltophilia* increased *P. aeruginosa* resistance to colistin.                | Ryan et al., 2008   |
| Essential oils of citronella and lemon | *Escherichia coli*, *Staphylococcus aureus*        | *S. aureus* and *E. coli* increased resistance to citronella and lemon essential oils when co-cultured. | Millezi et al., 2012 |
| Gentamicin      | *Enterococcus faecalis*, *Finegoldia magna*, *Pseudomonas aeruginosa*, *Staphylococcus aureus* | *P. aeruginosa* increased resistance to gentamicin when co-cultured with *S. aureus*, *E. faecalis*, and *F. magna*. | Dalton et al., 2011 |
| Hydrogen peroxide | *Pseudomonas aeruginosa*, *Staphylococcus aureus*       | *P. aeruginosa* induction of pigment synthesis and catalase upregulation in *S. aureus* increased its resistance to hydrogen peroxide. | Antonic et al., 2013 |
| Antibiotic            | Bacterial Strain(s)                                      | Mechanism                                                                 | Reference                                |
|-----------------------|----------------------------------------------------------|---------------------------------------------------------------------------|------------------------------------------|
| Ofloxacin             | *Candida albicans*, *Escherichia coli*                   | *C. albicans* β-1, 3-glucan, a matrix component, increased *E. coli* resistance to ofloxacin by acting as a barrier to its diffusion in the biofilm. | De Brucker et al., 2015                  |
| Ortho-phtalaldehyde acid | *Bacillus subtilis*, *Staphylococcus aureus*            | *B. subtilis* ypqP gene protected *S. aureus* from biocide action of ortho-phtalaldehyde acid. | Sanchez-Vizuete et al., 2015             |
| Peracetic acid        | *Bacillus subtilis*, *Staphylococcus aureus*            | *B. subtilis* ypqP gene protected *S. aureus* from biocide action of peracetic acid acid.. | Sanchez-Vizuete et al., 2015             |
| Polymyxin B           | *Pseudomonas aeruginosa*, *Staphylococcus aureus*       | *P. aeruginosa* induction of pigment synthesis and catalase upregulation in *S. aureus* increased its resistance to polymyxin B. | Antonic et al., 2013                    |
| Rifampicin            | *Dolosigranulum pigrum*, *Inquilinus limosus*, *Pseudomonas aeruginosa* | *P. aeruginosa* increased *I. limosus* and *D. pigrum* resistance to rifampicin. | Lopes et al., 2012                      |
| Sodium dodecyl sulphate (SDS) | *Pseudomonas fluorescens*, *Klebsiella pneumoniae*, *Pseudomonas aeruginosa* | *P. fluorescens* increased resistance to SDS when co-cultured with *K. pneumoniae* and *P. aeruginosa*. | Lee et al., 2014                        |
| Tobramycin            | *Pseudomonas aeruginosa*, *Stenotrophomonas maltophilia* | *P. aeruginosa* alginate protected *S. maltophilia* from tobramycin treatment- | Pompilio et al., 2015                   |
| Trimethoprim - sulfamethoxazole | *Haemophilus influenzae*, *Moraxella catarrhalis*  | *H. influenzae* signalling molecule AI-2 induced *M. catarrhalis* resistance to trimethoprim – sulfamethoxazole. | (Armbruster et al. 2010)                |
| Vancomycin            | *P. aeruginosa*, *S. aureus*                            | *P. aeruginosa* HQNO increases *S. aureus* resistance to vancomycin.      | Orazi and O'Toole, 2017                 |
Table 3. Examples of anti-biofilm molecules and their mechanisms of action, according to different strategies.

| Anti-biofilm strategy                                      | Mechanism of action                  | Molecules                                                                 | References                                                                 |
|-----------------------------------------------------------|--------------------------------------|---------------------------------------------------------------------------|---------------------------------------------------------------------------|
| Inhibition of bacterial attachment to surfaces             | Anti-adhesive surface properties      | Hydrophilic polymers and surfaces with microporous and nanoscale topography | Huang et al., 2002; Hsu et al., 2013; Kang et al., 2016                   |
| Antimicrobial surface properties                           | Anti-adhesive surface properties      | Antimicrobial peptides, essential oils, metallic nanoparticles, bacteriophages | Agnihotri et al., 2013; Wang et al., 2016; Zaltsman et al., 2017; He et al., 2018 |
| Interference with signal molecules modulating biofilm formation | AI degradation                      | Lactonases, acylases and oxidoreductases                                  | Wang et al., 2004; Lade et al., 2014; Chan et al., 2015                   |
| AISynthesis inhibition                                     | AISynthesis inhibition                | Halogenated furanone compounds, quercetin, cycloleucine, nickel and cadmium | Kim et al., 2008; Vega et al., 2014; Yadav et al., 2014; Gopu et al., 2015 |
| Al antagonization                                          | AIB antagonization                   | AHL analogues (cyclic sulfur compounds, phenolic compounds), AI-2 analogues (ursolic acid, isobutyl-4,5-dihydroxy-2,3-pentanediol (isobutyl-DPD) and phenyl-DPD), AIP analogues (cyclic peptides, RNA III) | Brackman and Coenye, 2015; Hossain et al., 2017                           |
| C-di-GMP signalling system inhibition                      | C-di-GMP signalling system inhibition | LP 3134, LP 3145, LP 4010, LP 1062                                       | Bachovchin et al., 2009; Sambanthamoorthy et al., 2014                    |
| Disruption of biofilm architecture                         | EPS/matrix degradation                | Polysaccharide-degrading enzymes (dispersin B, endolysins), nucleases (DNase I), proteases (proteinase K, trypsin) | Kaplan et al., 2003; Sugimoto et al., 2018                                |
| Biofilm dispersion                                         | Biofilm dispersion                    | Nitric oxide, cis-2-decenoic acid (CDA), EDTA, lactoferrin               | Banin et al., 2005, 2006; Barraud et al., 2009; Kumar Shukla and Rao, 2013; Marques et al., 2015 |