

REVIEWS

Shifting the balance: antibiotic effects on host–microbiota mutualism

Benjamin P. Willing*[§], Shannon L. Russell*^{†§} and B. Brett Finlay*[‡]

Abstract | Antibiotics have been used effectively as a means to treat bacterial infections in humans and animals for over half a century. However, through their use, lasting alterations are being made to a mutualistic relationship that has taken millennia to evolve: the relationship between the host and its microbiota. Host–microbiota interactions are dynamic; therefore, changes in the microbiota as a consequence of antibiotic treatment can result in the dysregulation of host immune homeostasis and an increased susceptibility to disease. A better understanding of both the changes in the microbiota as a result of antibiotic treatment and the consequential changes in host immune homeostasis is imperative, so that these effects can be mitigated.

Microbiota

The consortium of microorganisms that inhabit the body.

Mutualist

Pertaining to a member of the gut microbiota: a species that provides a service to the host while benefiting from the environment afforded by the host.

Extra-intestinal disease

A disease that manifests outside of the gastrointestinal tract but is influenced by the gut microbiota.

*Michael Smith Laboratories, The University of British Columbia.

[†]Department of Microbiology and Immunology, The University of British Columbia, Vancouver, British Columbia V6T 1Z4, Canada.

[§]These authors contributed equally to this work.

Correspondence to B.B.F.

e-mail:

bfinlay@interchange.ubc.ca
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The term *microbiota* refers to an abundant and diverse population of bacteria, archaea and fungi that reside in body sites that are exposed to the environment, with the highest concentrations of these organisms being found in the gastrointestinal tract. These *mutualists* contribute substantially to host phenotype, so extensively, in fact, that mammals have been described as ‘superorganisms’, and terms such as ‘ecological development’ have been coined to signify that development is a product of both host genetics and interactions with the microbiota¹. With recent advances in our understanding of the intestinal microbiota and its role in regulating immunity, we are becoming increasingly aware of the effects that antibiotics have on the resident microbial ecosystem and how they can adversely affect patient health. One of the most striking examples is antibiotic-associated diarrhoea (AAD) caused by the expansion of *Clostridium difficile*, and subsequent pseudomembranous colitis, which can result in death².

Interactions between microorganisms and the host direct the immune system locally and systemically³. Animal models demonstrate that antibiotic treatments affect both the innate and adaptive immune systems dynamically, affecting more than just immune development. Therefore, disturbances in a healthy adult microbiota can create opportunities for pathogens to infect and increase disease severity^{4–6}. Antibiotics also provide excellent tools to study the role of the microbiota in protecting from and contributing to intestinal and extra-intestinal diseases, including gastroenteritis induced by *Salmonella enterica* subsp. *enterica* serovar Typhimurium, multiple sclerosis, diabetes, autism and atopic diseases^{7–11} (FIG. 1; TABLE 1).

The microbiota protects the host from incoming pathogens through various mechanisms, including competition for space or binding sites, competition for nutrients by bacteria with similar nutrient requirements, and direct inhibition through the release of inhibitory molecules (reviewed in REF. 12). However, the focus of this Review is on how antibiotics affect the microbiota, how these changes affect immune homeostasis and how loss of immune homeostasis creates susceptibility to disease. Although the focus here is on negative impacts on the microbiota and the host, antibiotics will continue to be important tools in the battle against infectious disease. Therefore, strategies are required to allow mucosal and immunological integrity to be maintained when antibiotics are administered. New therapeutic approaches towards this aim include faecal bacteriotherapy from healthy donors following antibiotic treatment, and exogenous immune activation during the course of antibiotic therapy^{13,14}.

Impact of antibiotics on the microbiota

To understand the consequence of antibiotics on immune homeostasis, we must first understand how they affect the intestinal microbial ecosystem. Antibiotics are selectively administered to target a specific pathogenic population. However, most antibiotics produced have broad-spectrum activity so that they can be used to treat many diseases¹⁵. Therefore, although antibiotics are designed to target pathogenic organisms, related members of the microbiota are also affected, often leaving an imprint on the gut community long after the antibiotics have been removed^{16,17}. Antibiotic use also promotes the expansion of antibiotic-resistant strains, which can act as a lasting

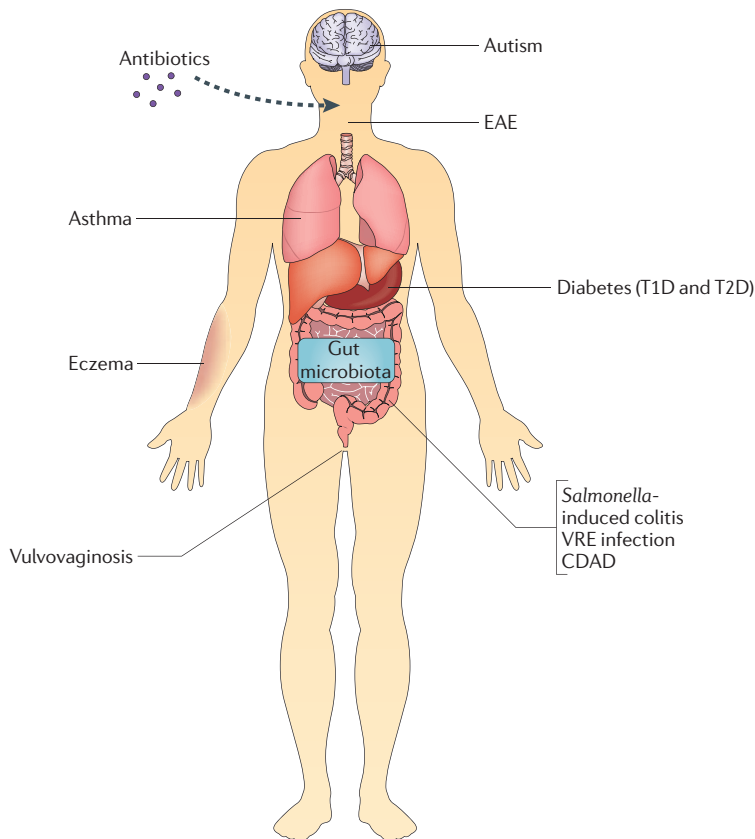


Figure 1 | Oral antibiotics have been associated with changes in numerous intra- and extra-intestinal diseases. Several associations between oral antibiotics and intra- or extra-intestinal diseases (indicated in the figure) have been made in clinical studies, whereas other such associations have only been made in experimental models to date. CDAD, *Clostridium difficile*-associated diarrhoea; EAE, experimental autoimmune encephalopathy; T1D, type 1 diabetes; T2D, type 2 diabetes; VRE, vancomycin-resistant *Enterococcus*.

antibiotics may target specific bacteria, but other species that exchange secondary metabolites or waste products with the targeted species are indirectly affected (FIG. 2). This phenomenon was seen for treatment with vancomycin, after which some Gram-negative populations were dramatically depleted, despite the antimicrobial activity of the antibiotic being restricted to Gram-positive organisms²³.

Antibiotics shift the composition of the microbiota in different ways depending on their spectrum of activity, a factor that can be useful in identifying which bacterial groups correlate to disease susceptibility or changes in host immunity. These antimicrobials also have different effects on diversity, bacterial biomass and the ability of the microbiota to recover post-treatment²⁴. Although the spectrum of the antibiotic is paramount, the administered dose is also extremely important for determining the ecological impact of a drug on the microbiota. Sub-therapeutic doses of antibiotics are used in the agriculture industry both prophylactically and to promote animal growth. Antibiotic growth promoters have been criticized because, although they do not reduce the total bacterial biomass, they shift the composition of the microbiota and promote dangerous levels of antibiotic resistance²⁵. In piglets that are given amoxicillin 1 day after birth, the microbiota remains substantially altered 5 weeks after treatment, further emphasizing the long-term effects of antibiotics on gut homeostasis²⁶. The therapeutic doses that are used in clinical practice are designed to minimize the effects on the host while maximizing pathogen clearance. These doses do not typically reduce the total bacterial biomass in the gut, but the elimination of subsets of the microbiota can shift the community in ways that promote colonization by opportunistic pathogens such as *C. difficile*²⁷ and *Candida albicans*²⁸. It should be noted that effects on the intestinal microbiota and mucosal immunity are not limited to orally administered antibiotics²⁹. Systemically delivered antibiotics can have impacts on the intestinal microbiota, as they can reach the gut through the biliary system³⁰. Even antibiotics that are secreted by the biliary system in only low amounts have been shown to have effects on the gut microbiota³¹. Antibiotic doses above therapeutic levels are used primarily for research purposes in which the depletion of the total bacterial biomass is required to test for a role of the microbiota.

reservoir for resistance genes in the gut microenvironment¹⁸. Antibiotic treatment is typically followed by a decrease in the diversity of the microbiota¹⁷, but the community is resilient and resembles the pretreatment state in a matter of days or weeks^{19,20}. Although most of the microbiota return to pretreatment levels, some members do not and are lost from the community indefinitely^{16,17}.

The microorganisms that are affected by antibiotic therapy are not limited to those that are directly targeted by the antibiotic. A complex network of co-dependence exists among members of the microbiota, driven by differential metabolite production and utilization pathways. These ecological interactions have been demonstrated in colonization models involving prominent members of the intestinal microbiota²¹. Microorganisms can be dependent on other colonizers for the provision of nutrients or secondary metabolites²², or for the removal of waste products that become toxic if they build up. For example, because of their co-dependence, the archaeon *Methanobrevibacter smithii* and the bacterium *Bacteroides thetaiotaomicron* were able to colonize germ-free mice more effectively when they were introduced together than when they were introduced alone²¹. Therefore,

Why antibiotics affect the host

Through the course of mammalian evolution, the microbiota has been selected to provide a vast number of services to the host, ranging from the provision of nutrients to the regulation of immune function. Host-microorganism interactions are surprisingly specific, as demonstrated by the inability of microorganisms from the human gut to appropriately regulate immune development of the mouse^{32,33}. These specific interactions, which are highly co-evolved processes, are altered or lost as a consequence of antibiotic treatment, leading to effects on host health. Changes that are induced by antibiotic treatment and that are important to microbial regulation of host immunity include the loss of bacterial

Bacteriotherapy

The introduction of the microbiota from a healthy donor into a patient with an imbalanced microbiota.

Sub-therapeutic

Below the dosage levels used to treat diseases.

Gut homeostasis

The dichotomy between preventing infection by luminal microorganisms and avoiding overt intestinal inflammation.

Co-evolved

Resulting from the evolution of interdependent species, such that each organism adapts to changes in the other.

Table 1 | Associations between antibiotics and disease phenotypes

Antibiotics	Clinical or experimental disease	Disease or disease model	Effects on the microbiota	Outcome of disease or disease model	Refs
Many antibiotics (e.g. clindamycin)	Clinical	CDAD	Uncharacterized	CDAD	118
Unspecified*	Clinical	Asthma or wheezing, eczema and allergic sensitization	Uncharacterized	Increased susceptibility to asthma or wheezing, but not to eczema, and no allergic sensitization	91–93
Cephalosporins	Clinical	Asthma	Suppression of lactobacilli and bifidobacteria, and overgrowth of <i>C. difficile</i>	Increased susceptibility to asthma	119
Unspecified*	Clinical	Vulvovaginal candidiasis	Loss of vaginal lactobacilli and overgrowth of <i>C. albicans</i>	Vulvovaginal candidiasis	28
Cocktail of kanamycin, gentamicin, colistin, metronidazole and vancomycin, and clindamycin injection	Experimental	CDAD	Antibiotic cocktail and clindamycin injections are required to achieve sufficient disruption of the microbiota prior to infection with <i>C. difficile</i>	Close resemblance to the human disease characteristics	77
High-dose streptomycin	Experimental	<i>S. Typhimurium</i> -induced colitis	95% of the total bacteria are eliminated	<i>S. Typhimurium</i> is better able to colonize the intestine of mice, creating a better model of gastroenteritis	82
Vancomycin or streptomycin	Experimental	<i>S. Typhimurium</i> -induced colitis	Shifts in composition occur (at the phylum level), but the total number of bacteria is unaffected	Treated hosts are rendered more susceptible to <i>S. Typhimurium</i> infection than untreated hosts, and both antibiotics allow better <i>S. Typhimurium</i> colonization	10
Streptomycin	Experimental	Cholera	Total number of bacteria is significantly reduced	<i>V. cholerae</i> is better able to colonize the intestine of guinea pigs, creating a better model to study cholera	120
Cefoperazone	Experimental	Allergic-airways disease	Total number of bacteria is significantly reduced, allowing overgrowth of <i>C. albicans</i>	Allergic-airways disease develops after challenge with mould spores	11, 95
Kanamycin	Experimental	Atopic dermatitis	Total number of anaerobes (specifically, lactobacilli) is significantly reduced	Clinical scores for atopic dermatitis are significantly higher	96
Cocktail of ampicillin, vancomycin, neomycin sulphate and metronidazole	Experimental	EAE	Total number of bacteria is significantly reduced	EAE clinical scores are significantly lower	9, 97, 98

C. albicans, *Candida albicans*; CDAD, *Clostridium difficile*-associated diarrhoea; EAE, experimental autoimmune encephalitis; *S. Typhimurium*, *Salmonella enterica* subsp. *enterica* serovar *Typhimurium*; *V. cholerae*, *Vibrio cholerae*. *No specific classes of antibiotic were identified.

ligands that are recognized by the host, alterations in the metabolites produced by the microbiota and the loss of specific bacterial signals.

Loss of bacterial ligands that are important for immune induction. An important mechanism through which the microbiota affects host immunity is the activation of microorganism-associated molecular pattern (MAMP) recognition receptors, including Toll-like receptors (TLRs) and NOD-like receptors (NLRs). These pattern recognition receptors (PRRs) are activated by ligands from intestinal microorganisms, including lipopolysaccharide (LPS), lipoteichoic acid (LTA), flagellin, CpG DNA and peptidoglycan³⁴. PRR signalling has been shown to be important in a range of homeostatic mechanisms that involve epithelial integrity and repair, as well as mucosal immune functions (reviewed in REF. 35). Depletion of

the microbiota using antibiotics certainly reduces the amount of MAMPs coming into contact with the epithelium; antibiotics targeting Gram-negative bacteria would primarily affect TLR4 and nucleotide-binding oligomerization domain-containing protein 1 (NOD1) activation, whereas antibiotics targeting Gram-positive bacteria would primarily affect TLR2 and NOD2 activation. Of note, not all LPS or LTA is equal in its ability to stimulate host responses³⁶. A number of studies have shown that depletion of the microbiota as a consequence of antibiotic treatment results in reduced TLR signalling and downstream regulation of innate defences^{5,37}.

Changes in bacterial metabolites. As well as having different abilities in terms of activating PRRs, each population of bacteria carries out specific metabolic processes. Correspondingly, the composition of the

Microorganism-associated molecular pattern
A molecule that is found as part of a microorganism and is recognized by cells of the host innate immune system.

Pattern recognition receptor
A protein that is expressed as part of the innate immune system and recognizes a microorganism-associated molecular pattern; when activated, these receptors elicit a downstream immune response.

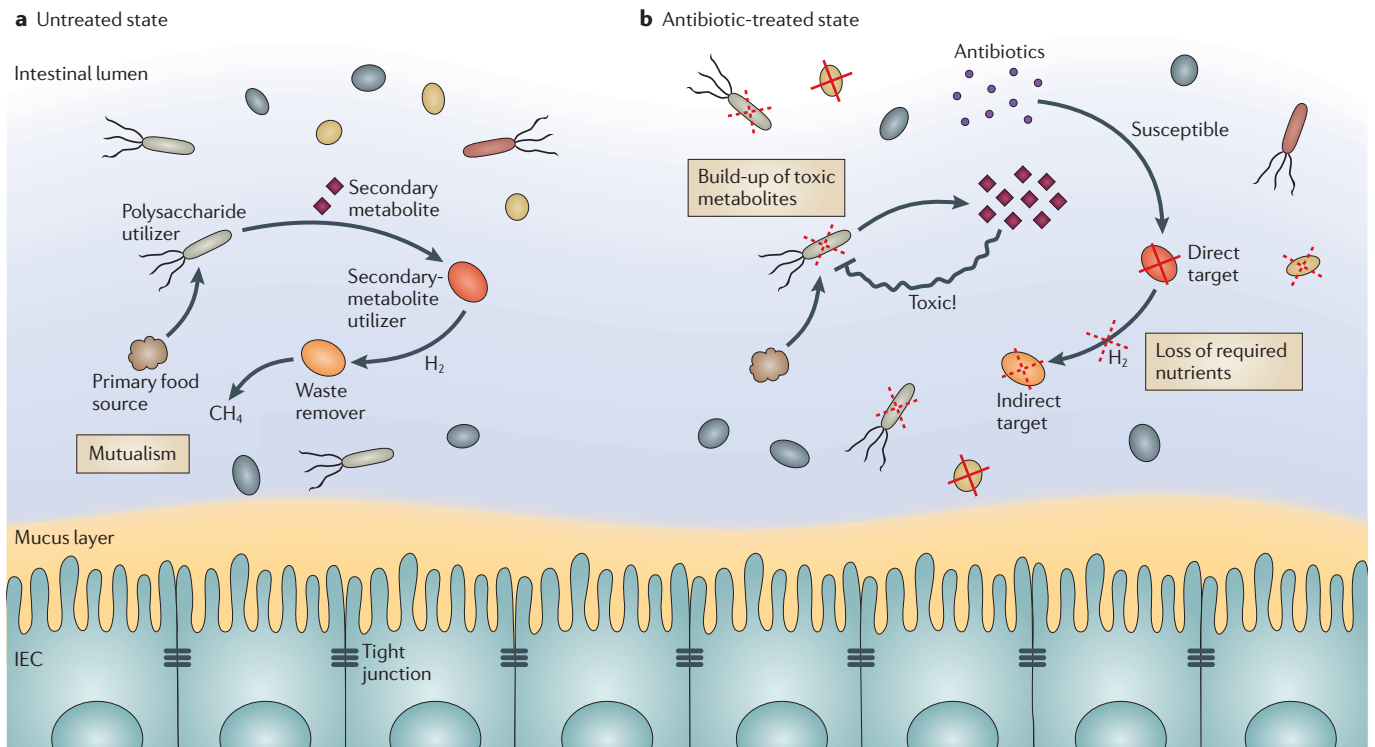


Figure 2 | Antibiotics have direct and indirect effects on the gut microbiota. a | Mutualism exists between gut symbionts and the host in the absence of antibiotics. Co-dependence among symbionts is exemplified by the relationships shown in the intestinal lumen in the figure. A polysaccharide utilizer acquires a primary food source from the environment, obtains energy through the metabolism of that food source and produces secondary metabolites as by-products into the surrounding environment. These secondary metabolites are used as fuel by neighbouring symbionts (secondary-metabolite utilizers) that are unable to break down complex primary food sources. The secondary-metabolite utilizers produce waste products, such as H₂, that are taken up by waste removers, which metabolize these waste products to make their own energy (for example, through methanogenesis). **b** | When antibiotics are introduced, mutualistic traits of the symbionts that are directly targeted (solid crosses) by the antibiotic are eliminated, and the microorganisms that are involved in those mutualistic associations are indirectly affected (dashed crosses). As an example, the loss of a targeted secondary-metabolite utilizer could promote the build-up of toxic secondary metabolites and/or eliminate a nutrient source for a waste remover. IEC, intestinal epithelial cell.

microbiota has been shown to correlate with both faecal and urinary metabolites^{38,39}. Thus, antibiotic treatment results in changes in the metabolic profiles of intestinal contents and faeces, as well as those of the blood and urine (following absorption of metabolites from the gut). Metabolic profiles have been studied in mice under various antibiotic regimens^{40,41}. A common feature of the metabolic profiles of antibiotic-treated mice and humans is a reduced production of short-chain fatty acids (SCFAs)^{40–43}. Two independent metabolomic studies of enrofloxacin- or vancomycin-treated mice showed decreased concentrations of amino acids and SCFAs in faeces from treated mice compared with concentrations in faeces from untreated mice, whereas oligosaccharide levels were increased in faeces from treated mice, indicating that microbial fermentation of carbohydrates was disrupted⁴¹. Although vancomycin had a dramatic effect on *Clostridium leptum* and *Clostridium coccooides* clusters (important butyrate-producing bacteria), these populations recovered 2–3 days post-treatment, coinciding with a recovery of fatty acid levels by day 7. In agreement with observations that some populations

of bacteria do not recover after antibiotic treatment, levels of some metabolites had not recovered weeks after the cessation of antibiotic treatment⁴¹.

The depletion of SCFAs following antibiotic treatment is important with regard to intestinal health and immunity. SCFAs are rapidly absorbed in the colon, providing a preferred energy source for colonocytes and regulating cell proliferation, differentiation, growth and apoptosis^{44–46}. Butyrate is the SCFA with the strongest effect on the cell cycle and has also been implicated in regulating many aspects of intestinal immunity. Effects on mucosal defence include improved tight junction assembly, antimicrobial secretion and mucin expression^{47–49}. Most studies have described butyrate as anti-inflammatory in the gut; for example, it has been shown to inhibit the constitutive expression of monocyte chemoattractant protein 1 (MCP1; also known as CCL2) in a dose-dependent manner, both in the absence and in the presence of LPS⁵⁰, and to reduce neutrophil production of reactive oxygen species⁵¹. These anti-inflammatory properties are generally considered to be beneficial, preventing overt inflammation,

Short-chain fatty acid
A fatty acid that is produced by the microbiota through the fermentation of host dietary fibre.

Metabolomic
Pertaining to the study of the collective metabolites in an environment.

oxidative stress and carcinogenesis⁵². Butyrate's anti-inflammatory effects on neutrophils may contribute to the increased infiltration of neutrophils that is observed in mice that are treated with streptomycin and then infected with *S. Typhimurium*, and could result in an increased inflammatory state in neutrophil-associated diseases. One mechanism by which butyrate acts is to induce peripheral blood mononuclear cells to produce prostaglandin E₂, a prostanoid that reduces the LPS-mediated induction of tumour necrosis factor (TNF) expression⁵³. Intriguingly, reduced concentrations of prostaglandin E₂ metabolites have been observed in patients with Crohn's disease, along with a disrupted microbiota, reduced butyrate production and uncontrolled intestinal inflammation^{39,54}. However, the effects of butyrate can be cell specific; therefore, caution should be used when interpreting the results of such studies. For example, the effect of butyrate on cathelicidin expression and antimicrobial activity is specific to colonocytes, as the molecule does not have the same effect on keratinocytes or monocytes⁴⁸. Butyrate has also been shown to increase the expression of NOD2, resulting in an increased induction of interleukin-8 (IL-8) following NOD2 activation by peptidoglycan in intestinal epithelial cells⁵⁵. As well as playing an important part in regulating mucosal immunity, SCFAs also regulate T cell energy, identifying these cells as targets for systemic and obesity-associated inflammation⁵⁶. Taken together, the data indicate an important anti-inflammatory role for butyrate, suggesting that the reduced butyrate levels seen during antibiotic treatment may contribute to increased inflammation following pathogen insult.

Butyrate is certainly not the only microbial metabolite that affects the host. However, in general, the metabolites produced by the microbiota are poorly understood. In a study examining metabolites in the faeces of patients with Crohn's disease, most (~90%) of the metabolites that correlated with disease could not be assigned to any metabolic pathway, indicating that the established databases are incomplete³⁹. Many of these unknown metabolites probably have important roles in regulating host physiology and mucosal immune homeostasis.

Loss of specific bacterial signals. The loss of specific bacteria also means the loss of specific signals that are associated with those bacteria. For example, although the process is not yet well understood, the population of segmented filamentous bacteria (SFB) is particularly adapted to induce the differentiation of T helper 17 (T_H17) cells⁶. Other specific microbial signals are involved in inhibiting inflammation. For example, *B. thetaiotaomicron* limited the inflammation induced by *Salmonella enterica* subsp. *enterica* serovar Enteritidis by activating nuclear export of nuclear factor-κB (NF-κB) through a peroxisome proliferator-activated receptor-γ (PPARγ)-dependent pathway⁵⁷. *Saccharomyces boulardii* reduced host inflammatory responses by secreting a heat-stable, water-soluble factor that blocked the degradation of NF-κB inhibitor-α (IκBa), thus reducing NF-κB binding to DNA and subsequent

IL-8 expression⁵⁸. Polysaccharide A, a component of *Bacteroides fragilis*, is able to induce FOXP3⁺ regulatory T cells to produce IL-10, preventing inflammation in an experimental model of infectious colitis⁵⁹. It is expected that many more microorganism-specific signals remain uncharacterized.

Antibiotic-related changes in immunity

The microbiota has many key roles in preparing the mucosal immune system for battle against infection. Germ-free animals have poorly developed lymphoid tissue, reduced and imbalanced T cell populations, increased intestinal permeability, and reduced expression of antimicrobial peptides (AMPs) and secretory immunoglobulin A⁶⁰. Although germ-free models have provided interesting contributions to our understanding of the role of the microbiota in immune development, these models do not address how changes in the microbiota later in the life of the host may affect host immunity. Studies using antibiotics have revealed that the microbiota not only contributes to development, but also is required for maintenance of the immune system (TABLE 2).

Many studies report reduced expression and secretion of AMPs on antibiotic treatment. AMPs, including C-type lectins, defensins and cathelicidins, are the first line of defence to incoming pathogens, as they are secreted into and maintained in the mucus layer⁶¹. AMPs have direct antimicrobial activity and are variably expressed between intestinal cell types; however, enterocytes, goblet cells and particularly Paneth cells all contribute to their production. Treatment with a combination of vancomycin, neomycin and metronidazole resulted in reduced expression of the gene encoding REG3γ, a C-type lectin that targets Gram-positive bacteria⁶². The expression of this gene could be corrected by administering LPS from Gram-negative bacteria, but not LTA from Gram-positive bacteria⁵. Conversely, treatment of mice with streptomycin has been shown to affect transcription of the mRNA encoding REG3β³⁷. The depletion of this transcript, and of the protein itself, seemed to be caused by reduced microbial activation of TLR2, as adding a TLR2 agonist resulted in recovery of expression³⁷.

Global gene analysis revealed that amoxicillin treatment also resulted in decreased expression of genes involved in antimicrobial defence, including those encoding α-defensins, matrilysin and phospholipase A2 (REF. 63). Although treatment with amoxicillin had no effect on total bacterial numbers in the small intestine, it nearly eradicated the lactobacilli. It should be noted that host responses differ depending on the location examined. Of the hundreds of genes affected by amoxicillin treatment, very few showed overlapping changes in the proximal small intestine, distal small intestine and large intestine. The only genes that were globally downregulated were those involved in antigen presentation, including those encoding major histocompatibility complex (MHC) class II and MHC class Ib complexes. Importantly, MHC class II complexes are required for tolerization to luminal antigens⁶⁴.

Energy

Reduced responsiveness to an antigen.

Antimicrobial peptide

A peptide that is secreted as part of the innate immune response and has broad-spectrum antibiotic activity.

Tolerization

The development of tolerance to innocuous antigens, including food and resident microbiota, thus contributing to intestinal homeostasis.

Table 2 | Antibiotic-associated changes in immunity

Antibiotic	Effects on the microbiota	Effects on immunity	Refs
Vancomycin, neomycin and metronidazole	• Reduced total bacteria	• Reduced expression of REG3 γ (recovered with LPS treatment)	5
Streptomycin	• Reduced total bacteria	• Reduced expression of REG3 β (recovered with a TLR2 agonist)	37
Amoxicillin	• Change in the microbial composition • Depletion of lactobacilli	• Reduced antimicrobial defence • Reduced expression of MHC molecules • Increased mast cell proteases	63
Polymyxin B	• Unknown	• Increased mast cell numbers	65
Vancomycin or ampicillin	• Depleted Gram-positive bacteria • Depleted SFB	• Depletion of T helper 17 cells	4,6
Ampicillin, gentamicin, metronidazole, neomycin and vancomycin	• Reduced total bacteria	• Reduced expression of IFN γ and IL-17A by CD4 $^+$ T cells • Reduced production of RELM β	69
Streptomycin and cefotaxime	• Reduced total bacteria	• Reduced expression of TLR2 and TLR4 in peritoneal macrophages • Macrophages less able to respond to LPS induction	70
Metronidazole	• Changes in the composition • More <i>Bifidobacterium</i> spp.	• Disrupted mucus layer • Increased colonization of intestinal crypts	75
Ampicillin, vancomycin, neomycin and metronidazole	• Reduced bacterial ligand CpG DNA	• Suppressed immune response to oral vaccination and infection	84
Amoxicillin and clavulanic acid	• Undetermined change in the microbiota	• Reduced systemic IgG	85
Ampicillin, neomycin sulphate, metronidazole and vancomycin	• Reduced total bacteria	• Reduced bacteriocidal activity of bone marrow-derived neutrophils	3

IFN γ , interferon- γ ; IgG, immunoglobulin G; IL-17A, interleukin-17A; LPS, lipopolysaccharide; MHC, major histocompatibility complex; RELM β , resistin-like molecule- β ; SFB, segmented filamentous bacteria; TLR, Toll-like receptor.

Although amoxicillin treatment reduced the expression of genes related to antigen presentation and the secretion of defence molecules, this treatment increased the expression of mast cell proteases. In fact, increased numbers of mucosal mast cells have been associated with the administration of other antibiotics⁶⁵. Mast cells play a part in host defence against pathogens⁶⁶, but they also have a key role in chronic intestinal inflammation⁶⁷. Treatment with antibiotics alone did not result in increased signs of inflammation; however, increased mast cell number and activity may predispose the host to exacerbated inflammatory reactions following insult with a pathogen.

Depleting specific subsets of the microbial population with antibiotics has provided clues as to which bacteria are responsible for regulating T cell differentiation and activation. The treatment of mice with vancomycin or ampicillin, which target Gram-positive bacteria, resulted in a depletion of T_H17 cells in the intestine, whereas treatment with metronidazole and neomycin sulphate, which target strict anaerobes and Gram-negative bacteria, respectively, had no effect on the T_H17 cell population⁴. It was this preliminary evidence that identified Gram-positive bacteria, and more recently SFB, as major contributors to this T cell phenotype⁶. It has also been shown that specific populations of mutualistic bacteria — not including those obligate anaerobes that are affected by metronidazole treatment — are required for maintaining IL-17 production by $\gamma\delta$ T cells⁶⁸.

The reduced frequency of CD4 $^+$ T cells expressing interferon- γ (IFN γ) or IL-17A following a 10-day antibiotic treatment period (using a combination of ampicillin,

gentamicin, metronidazole, neomycin and vancomycin) supports the role of continued microbial signalling in maintaining effector T cell populations in the gut⁶⁹. As well as affecting T cell activity, this combination of antibiotics was found to reduce the production of resistin-like molecule- β (RELM β), another important AMP produced by goblet cells⁶⁹. The changes that were seen in the microbial population with this antibiotic treatment were extensive, but in general the species most strongly affected were members of the phylum Firmicutes, whereas certain members of the phyla Bacteroidetes and Proteobacteria were able to persist⁶⁹.

As well as influencing antimicrobial defences, antibiotics can affect the activity of immune cells. The eradication of enteric bacteria from 8-week-old mice after 3 weeks of antibiotic administration has recently been shown to reduce the expression of TLR2 and TLR4 in peritoneal macrophages⁷⁰. This reduced expression of TLRs resulted in a markedly impaired response to challenge with LPS, indicating that continued mucosal stimulation by enteric bacteria is required for the priming of macrophages in preparation for invasion⁷⁰. How this phenomenon is affected by short-term antibiotic therapy or more clinical doses has yet to be shown.

Mucosal imbalance leads to susceptibility

Changes in immunity, such as those that occur as a consequence of antibiotic treatment, result in increased susceptibility to infection by pathogens (FIG. 3). Antimicrobial secretions have been shown to be very important for impeding colonization of enteric pathogens^{5,71}. The disruption of mucus layer integrity, as is seen in mucin 2

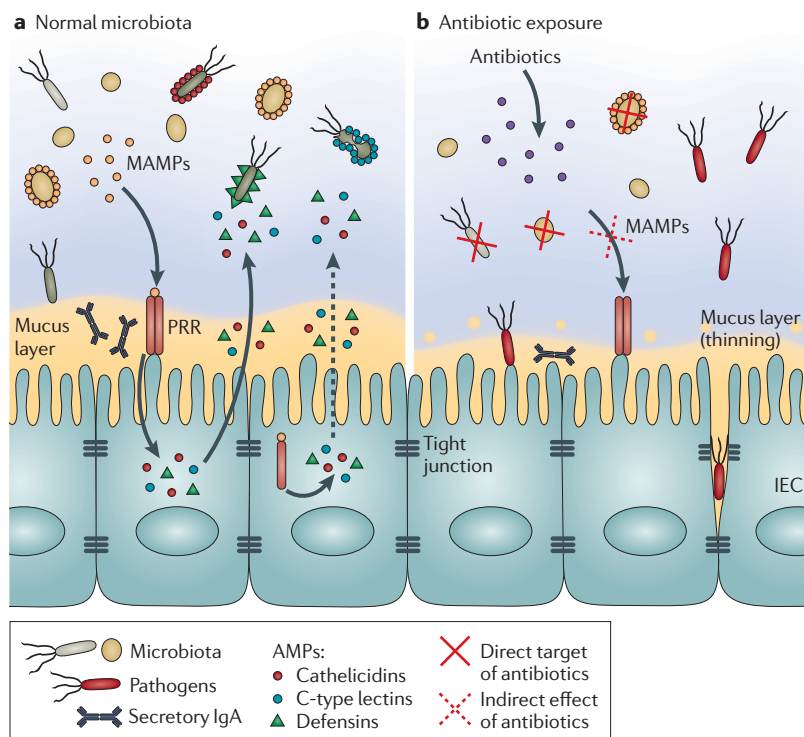


Figure 3 | Antibiotics alter epithelial homeostasis in the gut and enhance host susceptibility to incoming pathogens. **a** | In the absence of antibiotics, epithelial homeostasis is maintained. Microorganism-associated molecular patterns (MAMPs) from the microbiota are sensed intracellularly and extracellularly by pattern recognition receptors (PRRs) (that is, Toll-like receptors and NOD-like receptors). PRRs that recognize MAMPs from the microbiota instruct intestinal epithelial cells (IECs) called Paneth cells to secrete antimicrobial peptides (AMPs; for example, C-type lectins, defensins and cathelicidins) into the adjacent mucus layer. Secretory immunoglobulin A (IgA) and AMPs are important for controlling bacterial numbers and maintaining epithelial homeostasis. Tight junctions and a thick mucus layer prevent incoming pathogens from damaging the mucosal barrier. **b** | Antibiotic exposure disrupts epithelial homeostasis. Subsets of the normal microbiota are lost, thus reducing epithelial exposure to microbiota-derived MAMPs. Fewer MAMPs result in decreased host–microbiota signalling (that is, decreased PRR stimulation), which leads to decreased AMP production by IECs. Some antibiotics cause thinning of the mucus layer and disruption of tight junctions, rendering the host more susceptible to damage to the mucosal barrier, caused by incoming pathogens. The loss of AMP production and of the integrity of the epithelial cell barrier facilitates successful invasion by enteric pathogens.

(*Muc2*)-knockout mice, allows bacteria to contact the epithelium and, indeed, results in increased attachment of the enteric pathogen *Citrobacter rodentium*^{72,73}. Treatment with metronidazole — an antibiotic that targets anaerobic bacteria — affects the expression of MUC2, reduces the integrity of the mucous layer and accelerates mucosal attachment of *C. rodentium*⁷⁴. Intriguingly, streptomycin treatment did not result in reduced MUC2 expression, thus implying that bacteria which are obligate anaerobes (which are targeted by metronidazole but not streptomycin) are involved in triggering MUC2 expression. Treatment of rats with metronidazole also resulted in modifications in the mucus layer, resulting in greater microbial colonization of intestinal crypts⁷⁵.

Vancomycin-resistant *Enterococcus* (VRE) was also able to colonize more effectively in mice treated with antibiotics. Increased colonization was not a result of reduced

competition, but was instead due to a reduced production of the antimicrobial peptide REG3γ, as a consequence of depleted microbial signals⁵. A recent study of patients undergoing allogeneic haematopoietic stem cell transplantation also revealed that antibiotic treatment enabled VRE strains to dominate the microbiota, an event shown to precede bacterial invasion into the bloodstream⁷⁶. It is likely that parallels can be drawn between infection with VRE and with *C. difficile*. Antibiotic-induced susceptibility to *C. difficile* has been reproduced in a mouse model, providing a means of identifying how changes in mucosal immunity correlate with disease⁷⁷. However, it has yet to be determined whether host resistance to colonization by *C. difficile* is a product of changes in host immunity, or due to direct inhibition or competition from the microbiota. Certainly, immunological changes would be expected, as demonstrated in certain cases of AAD: changes in the composition of the microbiota in individuals with AAD have revealed a marked decrease in the prevalence of butyrate-producing bacteria⁷⁸.

It has recently been shown that *S. Typhimurium* uses gut inflammation to its competitive advantage when colonizing the gut⁷⁹. Specifically, reactive oxygen species produced on initial pathogen insult react with luminal sulphur compounds to form tetrathionate, a respiratory electron acceptor that is used by *S. Typhimurium* to gain a competitive advantage⁸⁰. Interestingly, the inflammation that is activated on initial infection and required for *S. Typhimurium* to expand and cause colitis does not occur in mice unless they are pretreated with antibiotics¹⁰. Probiotics have been shown to prevent colitis by inhibiting P-selectin upregulation and, thus, reducing the interactions between endothelial cells and both leukocytes and platelets⁸¹. Therefore, although this has not yet been shown experimentally, one might speculate that the loss of microbiota-produced anti-inflammatory signals that is known to occur on antibiotic treatment (as discussed above for butyrate) is involved in the increased inflammation that is seen on infection with *S. Typhimurium*. The high-dose streptomycin pretreatment model used in *S. Typhimurium*-induced colitis suggests that the microbiota as a whole was inhibiting *S. Typhimurium* colonization in the gut⁸². However, low-dose administration of streptomycin and vancomycin also enabled colonization by *S. Typhimurium*¹⁰, suggesting that different subsets of the microbiota are involved in modulating susceptibility to *S. Typhimurium* infection in the intestine. High-dose streptomycin treatment was also found to cause alterations in the microbiota that persisted for several weeks and resulted in continued susceptibility to *S. Typhimurium*-induced gastroenteritis⁸³.

Antibiotic-mediated depletion of bacteria has also been shown to affect the development of an adaptive immune response. Reducing the amount of CpG DNA (a bacterial MAMP) resulted in suppressed immune responses to oral vaccination with an antigen and to oral infection with a protozoan parasite⁸⁴. Combined treatment with amoxicillin and cluvulanic acid also resulted in reduced serum IgG levels, although IgA concentrations were unaffected⁸⁵, suggesting that systemic effects occurred.

Systemic effects

We are now learning that germ-free mice also have several extra-intestinal deficits, indicating that the effects of the microbiota on the host — which were previously believed to be restricted to the gut — have systemic implications. Antibiotic-induced perturbation of the microbiota demonstrated that NOD1-mediated recognition of peptidoglycan from Gram-negative bacteria is required for the generation of adaptive lymphoid tissues⁸⁶. Microbiota-derived peptidoglycan has been shown to interact with NOD1 (REF. 3) and to therefore systemically prime the innate immune system. This priming occurs through a mechanism that involves peptidoglycan from the microbiota entering the circulation, binding NOD1 receptors in bone marrow-derived neutrophils and initiating a signalling cascade that enhances the ability of these neutrophils to kill microorganisms⁸⁷. The concept of the microbiota affecting systemic immunity is not new: over 10 years ago, antibiotics were used to show that antigens from the microbiota were involved in regulating cell-mediated responses by priming CD4⁺ T cells to secrete IL-4 (in other words, to become T_H2 polarized)⁸⁸. This was observed even after challenge with *Leishmania major*, a pathogen requiring a T_H1-type response for clearance.

Antibiotics have also been used to investigate the role of extra-intestinal diseases that are caused by gut-specific microorganisms. Over the past few decades, there has been an increase in the prevalence of immunological disorders (both of autoimmune and of atopic origin) in the developed world. These observations, in combination with causal relationships established by animal models and clinical studies, have provided increased support for the hygiene hypothesis^{89,90}. Recent epidemiological studies have associated early antibiotic use with an increased risk of atopy^{91–93}, encouraging an extension of the hygiene hypothesis, coined the ‘microflora hypothesis’. This is the theory that reduced microbial exposure during childhood inhibits normal maturation of the intestinal microbiota, altering immune development in a way that enhances the symptoms of allergic hypersensitivity⁹⁴. Although these studies remain correlative, antibiotic exposure in animal models of allergic-airways disease^{11,95} and atopic dermatitis⁹⁶ support these findings; however, most of the mechanisms involved in these models remain undefined.

Although no studies have linked antibiotic exposure to autoimmunity in a clinical setting, experiments in animal models have provided some exciting insights. Oral antibiotic treatment has demonstrated the substantial impact of the microbiota on experimental autoimmune encephalomyelitis (EAE), the experimental model of multiple sclerosis. Depleting the intestinal microbiota with a cocktail of antibiotics impaired the development of EAE. Protection was accompanied by diminished pro-inflammatory responses and an accumulation of FOXP3⁺ regulatory T cells in the peripheral lymph nodes⁹⁷. Interestingly, antibiotic-treated mice reconstituted with polysaccharide A-producing *B. fragilis* remained protected from EAE, whereas colonization of mice with a polysaccharide A-deficient mutant of *B. fragilis* restored EAE susceptibility. Recent data suggest that polysaccharide A enhances the conversion of naive

CD4⁺ T cells into IL-10-producing FOXP3⁺ regulatory T cells through induction by tolerogenic CD103⁺ DCs, an effect that protects mice against EAE induction^{9,98} (FIG. 4).

Antibiotic studies have helped to identify a role for the intestinal microbiota in several complex multifactorial disorders, such as metabolic syndrome and autism. Interestingly, there is evidence to suggest that certain antibiotic regimens may actually improve insulin sensitivity in both animals^{8,99} and humans¹⁰⁰. Associations between extensive antibiotic use and late-onset autism (which develops at 18–24 months of age) suggest that a reassembly of the microbiota may promote the overgrowth of microorganisms that trigger autistic behavioural patterns¹⁰¹. According to one study, oral vancomycin treatment actually reduces autistic symptoms in some patients, suggesting that certain gut microorganisms may influence the onset of this disorder¹⁰².

Challenges

Many interactions between host and microorganism require a functional immune system; therefore, an antibiotic approach is, in some ways, better than a gnotobiotic approach for understanding host–microbial interactions, as immune function in gnotobiotic animals is drastically different from that in conventional animals. Although this is an important limitation of the gnotobiotic system, using antibiotics to understand host–microbial interactions has many of its own limitations.

One of the major challenges associated with using antibiotics to study host–microbial interactions is the direct effect that antibiotics can have on the immune system. Macrolide antibiotics in particular have been described as having many non-antimicrobial effects on host physiology¹⁰³. This class of antibiotics typically decreases the production of pro-inflammatory cytokines, including IL-1 β , IL-8 and macrophage inflammatory protein, and decreases neutrophil infiltration^{104,105}. The aminoglycoside gentamicin has been shown to have direct inhibitory effects on the production of TNF and MCP1 in proximal tubular cells, despite an increased accumulation of the mRNAs encoding these proteins¹⁰⁶. Similarly, although not demonstrated as a direct effect in the gut, LPS induced greater transcription of TNF in mice treated with gentamicin. Concanamycin B has also been found to inhibit the acidification of endosomes and lysosomes in macrophages¹⁰⁷. Certainly, this could affect the ability of macrophages to clear intracellular pathogens.

Reproducing antibiotic-induced alterations in the microbiota is also an important challenge. How the microbiota responds to antibiotic administration is largely dependent on the length of exposure and the number of repeated exposures¹⁰⁸. It has been well documented that the number of antibiotic-resistance genes in the microbiota is dramatically increased following antibiotic exposure, causing a reorganization of the phylogenetic hierarchy of the community^{17,18}. It is thought that repeated exposure to the same antibiotic fails to reproduce the same compositional shifts owing to the overgrowth of these resistant strains. There is also considerable variability in the composition of the microbiota among individuals, particularly at the genus and species levels^{109,110}.

Hygiene hypothesis

The hypothesis that insufficient exposure to microorganisms, parasites and infectious agents alters natural immune development, leading to an increased susceptibility to allergic diseases.

Atopy

An allergic reaction that is characterized by immunoglobulin E-mediated inflammation.

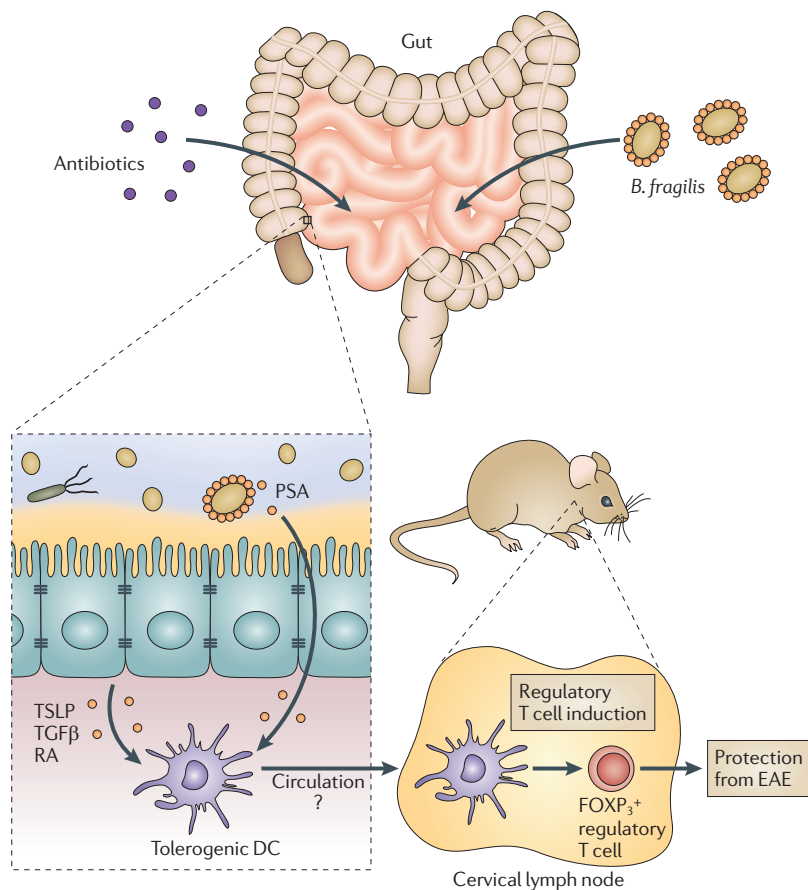


Figure 4 | Antibiotics used as tools to understand the role of polysaccharide A from *Bacillus fragilis* in protection against experimental autoimmune encephalopathy. Following antibiotic-mediated disruption of the intestinal microbiota, wild-type and polysaccharide A (PSA)-deficient *Bacteroides fragilis* were administered to mice. The wild-type bacteria protected against experimental autoimmune encephalomyelitis (EAE), but the PSA-deficient mutant did not⁹. The suggested mechanism of protection involves transport of the PSA antigen across the epithelial cell barrier, induction of tolerogenic dendritic cells (DCs) in the underlying lamina propria in the presence of epithelial cell-secreted cytokines, and subsequent transport of the tolerogenic DCs in the circulation (through an unknown mechanism) to cervical lymph nodes. In the cervical lymph nodes, tolerogenic DCs promote the induction of FOXP3⁺ regulatory T cells. The expansion of FOXP3⁺ regulatory T cells in the cervical lymph node corresponds with protection against demyelination of the central nervous system in EAE. TSLP, thymic stromal lymphopoietin; TGFB, transforming growth factor- β ; RA, retinoic acid.

As such, antibiotic treatment results in individualized responses to perturbation of the microbiota¹⁶. Similarly, different mouse strains¹¹, or even the same mouse strain housed in a different animal facility, respond differently to the same disease model.

Because even narrow-spectrum antibiotics do not target a single species or even a single genus, identifying those changes in the microbiota that are associated with a subsequent downstream effect (that is, an immune response, or an altered susceptibility to a pathogen or disease) is challenging. Many antibiotic studies remain correlative because no direct cause-and-effect relationship has been identified.

Transplantation is an alternative to antibiotic methods of microbiota manipulation. However, reintroducing specific organisms into a host is challenging owing to the

lack of cultured organisms available. Transfer models are being developed in which faeces or caecal contents can be used to inoculate a host with a particular community¹¹²; however, this involves the transfer of a complete microbiota rather than single organisms. Achieving efficient transfer is difficult, which is why many transplants are performed in germ-free mice. More effective, targeted approaches are needed to supplant the correlative data that has been accumulated using antibiotics.

Where do we go from here?

Antibiotics create stable, lasting alterations in the gut microbiota, weakening the fidelity of this community by disrupting the host–microorganism associations that have co-evolved to maximize mutual benefit. The loss of these mutualistic traits affects the remaining community members and their hosts across generations, shifting homeostatic mechanisms toward potentially unfavourable outcomes¹⁰⁸. Although this Review focuses on the role of antibiotics in disrupting the homeostatic balance of immunity, it must be noted that antibiotics can also restore a homeostatic state in cases in which the host is unable to regulate the microbiota. A lack of immunoglobulin genes in the host results in a 100-fold expansion of the microbiota, resulting in hyperplasia of lymphoid follicles; however, antibiotic treatment corrects this response¹¹³.

With an increasing number of studies demonstrating that antibiotics have detrimental effects on the immediate and long-term health of the host, there is a push to develop treatments that are better targeted to specific infecting pathogens; the challenges are not limited to the development of these drugs, but also include methods for rapid, accurate diagnoses in the clinic¹⁵. Because eliminating the need for antibiotics is unrealistic at this time, a better understanding of how antibiotics impact the microbiota and host immunity is needed, so that strategies to mitigate these effects can be developed. Bacteriotherapy is rapidly becoming a recognized treatment for antibiotic-induced *C. difficile*-associated diarrhoea¹³. As some microorganisms do not return in the microbiota after antibiotic therapy, it might be prudent to collect faecal samples before treatment, to be used as a source of ‘individualized’ healthy microorganisms that could be administered post-treatment. Moreover, probiotic intervention strategies are being implemented following antibiotic therapy in infants with necrotizing enterocolitis (NEC)¹¹⁴, as well as in individuals being treated for AAD¹¹⁵ and bacterial vaginosis¹¹⁶. Activation of TLR4 seems to be one of the mechanisms involved in bacteria-mediated protection from diseases, including NEC¹¹⁷, suggesting that TLR agonists could be administered during antibiotic therapy to help restore immune homeostasis. Similarly, systemic administration of flagellin has been used to experimentally limit infection with VRE; through activation of TLR5, systemically administered flagellin restored antibiotic-impaired immune defences (specifically, REG3 γ expression), drastically reducing colonization by VRE¹⁴. Although most of these approaches are still experimental, they hold substantial promise as adjunct therapies to antibiotics.

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Competing interests statement

The authors declare no competing financial interests.

FURTHER INFORMATION

B. Brett Finlay's homepage:
<http://www.finlaylab.msl.ubc.ca/index.html>
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