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The impact of intestinal inflammation on the nutritional environment of the gut microbiota

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Abstract

The intestinal epithelium is a single cell barrier separating a sterile mucosal tissue from a large microbial community dominated by obligate anaerobic bacteria, which inhabit the gut lumen. To maintain mucosal integrity, any breach in the epithelial barrier needs to be met with an inflammatory host response designed to repel microbial intruders from the tissue, protect the mucosal surface and repair injuries to the epithelium. In addition, inflammation induces mechanisms of nutritional immunity, which limit the availability of metals in the intestinal lumen, thereby imposing new selective forces on microbial growth. However, the inflammatory host response also has important side effects. A by-product of producing reactive oxygen and nitrogen species aimed at eradicating microbial intruders is the luminal generation of exogenous electron acceptors. The presence of these electron acceptors creates a new metabolic niche that is filled by facultative anaerobic bacteria. Here we review the changes in microbial nutrient utilization that accompany intestinal inflammation and the consequent changes in the composition of gut-associated microbial communities.

Keywords

Intestinal inflammation; microbiota; nutritional immunity; anaerobic respiration

Introduction

During intestinal inflammation the epithelium plays an important role in mounting responses that are aimed at clearing the mucosal surface from microbes. For example, production of IFN- γ during inflammation results in the activation of DUOX2 (dual function NADPH oxidase 2) [1], NOX1 (NADPH oxidase 1) [2] and iNOS (inducible nitric oxide synthase) [3] in epithelial cells. Reactive oxygen species (ROS) produced by DUOX2 and NOX1 and reactive nitrogen species (RNS) generated by iNOS create a hostile environment in close proximity to the mucosal surface. Furthermore, the pro-inflammatory cytokine interleukin

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(IL)-22 induces the luminal release of the antimicrobial proteins lipocalin-2, calprotectin, RegIII β (regenerating islet-derived 3 beta) and RegIII γ from epithelial cells [4-6].

These epithelial defenses can be augmented by the transmigration of neutrophils into the intestinal lumen as the severity of intestinal inflammation increases. Upon transmigration, the phagocyte NADPH oxidase (PHOX), superoxide dismutase (SOD) and myeloperoxidase (MPO) of neutrophils generate additional ROS in the gut lumen. Subsequent lysis of neutrophils in the intestinal lumen releases calprotectin, which constitutes approximately 40% of their cytoplasmic content [7]. As a result, neutrophils are the main sources of luminal calprotectin during severe intestinal inflammation [8].

Some of the antimicrobials released into the intestinal lumen are bacteriocidal, thereby protecting the mucosa from infection. For instance, release of the C-type lectin RegIII γ contributes to luminal clearance of opportunistic pathogens, such as *Listeria monocytogenes* or vancomycin-resistant *Enterococcus faecium*, which are both members of the class Bacilli within the phylum Firmicutes [9,10]. Chronic granulomatous disease, an illness caused by PHOX-deficiency, illustrates that the generation of ROS by phagocytes is essential for preventing recurrent bacterial infections [11-13]. It is thus likely that upon transmigration into the lumen, the respiratory burst of neutrophils aids in clearing bacteria from the vicinity of the mucosal surface. However, recent evidence suggests that in addition to its bacteriocidal effects, the inflammatory host response has also a profound impact on the nutritional environment in the gut lumen, which can lead to alterations in the composition of gut-associated microbial communities (microbiota). Here we review these novel hypothesis and the underlying mechanisms.

Nutritional immunity changes the rules for microbial contestants

One subset of antimicrobial proteins released into the intestinal lumen during inflammation functions in limiting the availability of trace elements required for bacterial growth, such as iron and zinc, a host defense mechanism known as nutritional immunity. Bacteria acquire ferric iron (Fe³⁺) by releasing high-affinity iron chelators, termed siderophores (reviewed in [14]). Enterobactin, a cyclic trimer of N-(2,3-dihydroxybenzoyl)-L-serine, is the siderophore produced by most members of the Enterobacteriaceae, a family of facultative anaerobic bacteria belonging to the class Gammaproteobacteria within the phylum Proteobacteria [15-17]. After chelating iron, the Fe³⁺-enterobactin complex is transported actively by an energy-coupled outer membrane receptor protein into the periplasm. The energy required for transporting the Fe³⁺-enterobactin complex across the outer membrane is provided by the proton motive force of the cytoplasmic membrane, which is transmitted to the outer membrane via the TonB protein (reviewed in [14]).

Lipocalin-2 prevents bacterial iron acquisition by binding and sequestering enterobactin [18-20]. While uptake of Fe³⁺-enterobactin is a viable strategy for obtaining iron in the non-inflamed intestine, the epithelial release of lipocalin-2 during conditions of inflammation inhibits growth of bacteria relying solely on enterobactin for iron acquisition. Thus, bacteria acquiring iron through mechanisms that are not inhibited by lipocalin-2 gain a relative luminal growth advantage in the inflamed gut. This concept was first described in

Salmonella enterica, a member of the Enterobacteriaceae that secretes enterobactin along with a glycosylated derivative of enterobactin, termed salmochelin [21]. Salmochelin is not sequestered by lipocalin-2, thereby conferring resistance against this antimicrobial protein [22,23]. Deletion of the *iroN* gene, which encodes the TonB-dependent outer membrane siderophore receptor [24], renders *S. enterica* unable to utilize salmochelin [21]. As a result, an *S. enterica iroN* mutant solely relies on enterobactin for iron-acquisition. Compared to wild-type bacteria, growth of a *S. enterica iroN* mutant in the lumen of the mouse gut is reduced in the presence, but not in the absence of intestinal inflammation. Furthermore, *S. enterica* wild type and *iroN* mutant grow equally well in the inflamed gut of lipocalin-2-deficient mice [5]. Thus, luminal growth of lipocalin-2 resistant bacteria is favored in the inflamed gut, but not in the absence of intestinal inflammation.

A second metal that is sequestered by the host during inflammation through the release of antimicrobial proteins into the intestinal lumen is zinc. Calprotectin, a heterodimer composed of S100A8 and S100A9, inhibits bacterial growth in tissue by chelating both manganese and zinc [25]. Recent studies suggest that the transepithelial migration of neutrophils and the subsequent release of calprotectin from dead neutrophils reduce the availability of zinc in the intestinal lumen [8]. Zinc is transported across the cytoplasmic membrane of *S. enterica* by the high-affinity ABC (ATP binding cassette) transporter ZnuABC [26]. Compared to the *S. enterica* wild type, luminal growth of a *znuA* mutant is impaired in the inflamed intestine of wild type mice, but not in the inflamed intestine of S100A9-deficient mice [8]. These data support the idea that by overcoming the calprotectin-mediated host zinc sequestration, bacterial high-affinity zinc acquisition confers a luminal fitness advantage during colitis.

Above examples illustrate that the inflammatory host response can influence bacterial growth by changing the nutritional environment in the intestinal lumen. As a result, bacterial metal acquisition strategies that bestow no apparent growth benefit in the healthy gut can confer a luminal fitness advantage in the inflamed intestine. In other words, the host response can alter the contest rules that govern microbial competition for metals.

Interestingly, reducing the availability of metals brings microbes, which rely on similar iron acquisition strategies, into a contest. For example, the commensal *Escherichia coli* strain Nissle 1917, a member of the family Enterobacteriaceae, elaborates four siderophores, including enterobactin, salmochelin, aerobactin and yersiniabactin [27-29]. Of these siderophores, only enterobactin is sequestered by lipocalin-2. Co-colonization with *E. coli* Nissle 1917 reduces luminal growth of the pathogenic *S. enterica* in wild-type mice, but not in lipocalin-2-deficient mice. Furthermore, co-colonization of mice with a siderophore utilization-deficient *E. coli* Nissle 1917 *tonB* mutant does not reduce the ability of *S. enterica* to grow in the intestinal lumen [30]. These data suggest that by lowering the availability of iron in the lumen, the host inflammatory response can alter the outcome of a competition between bacterial species that utilize overlapping siderophore repertoires.

Microbial metabolism in the healthy large intestine

In addition to conferring nutritional immunity by lowering the availability of metals in the intestinal lumen, the host response changes the luminal environment by generating inflammation-derived nutrients as a by-product. The resulting bloom of bacterial species that can utilize inflammation-derived nutrients can alter the composition of gut-associated microbial communities. To understand how inflammation-derived nutrients alter the growth conditions in the large bowel, it is important to first grasp the nutrient acquisition and utilization strategies that characterize a balanced microbiota, which inhabits the healthy gut.

In healthy individuals, obligate anaerobic bacteria belonging to the classes Bacteroidia (phylum Bacteroidetes) and Clostridia (phylum Firmicutes) dominate microbial communities inhabiting the anaerobic environment of the lower gastrointestinal tract [31]. Since simple carbohydrates and proteins are digested and absorbed in the upper gastrointestinal tract, complex carbohydrates (e.g. fiber or mucus carbohydrates) or non-digestible proteins (e.g. gluten) are the main nutrients supporting growth of Bacteroidia and Clostridia in the large bowel. Oxygen or other exogenous electron acceptors are not available in the healthy distal gut to support respiration. Thus, microbes rely largely on fermentation of carbohydrates and amino acids to generate energy via substrate-level phosphorylation and to acquire carbon and nitrogen for the biosynthesis of proteins, carbohydrates, lipids and nucleotides.

To maintain redox balance during fermentation, electrons have to be transferred from NADH onto organic compounds, such as phosphoenolpyruvate, thereby generating metabolic end products that are released. Microbiota-derived fermentation end products that commonly accumulate in the gut lumen include formate, acetate, propionate, butyrate, lactate and hydrogen (H₂) (Fig. 1A). Some bacteria, such as *Bacteroides fragilis*, maintain redox balance by transferring electrons onto fumarate to generate succinate, a process known as fumarate respiration (reviewed in [32]). During this process, *B. fragilis* fixes host-derived carbon dioxide (CO₂) onto phosphoenolpyruvate to generate oxaloacetate, which is converted by reversing reaction of the tricarboxylic acid (TCA) cycle into the endogenous electron acceptor fumarate [33]. Succinate is released as a metabolic end product of fumarate respiration. Thus fumarate respiration and fermentation have in common that metabolically valuable phosphoenolpyruvate is removed from anabolic reactions and converted into metabolic end products to maintain redox balance.

Metabolic end products generated by Bacteroidia and Clostridia change the nutritional environment for both the host and other intestinal microbes. For example, some metabolic end products of Bacteroidia and Clostridia, such as butyrate, confer benefit to the host by providing nutrition for colonocytes, which mitochondrially oxidize this compound to carbon dioxide [34] (Fig. 1A). Other metabolites, such as hydrogen, are consumed by obligate anaerobic sulfate-reducing bacteria. Sulfate-reducing bacteria of the genus *Desulfovibrio* (class Gammaproteobacteria, family Desulfovibrionaceae) in turn generate the genotoxic gas hydrogen sulfide (H₂S) in the distal gut [35-37]. To avoid toxicity, host colonocytes express sulfide oxidases in their apical membrane that oxidize hydrogen sulfide to generate harmless thiosulfate (S₂O₃²⁻) [38,39].

By-products of the host response support anaerobic respiration

Inflammation causes a dramatic change in the nutritional environment in the large intestine by generating exogenous electron acceptors. This concept was first established by showing that a by-product of neutrophils mounting a respiratory burst in the gut lumen is the oxidation of thiosulfate to tetrathionate ($S_4O_6^{2-}$) [40] (Fig. 1B). *S. enterica* can use tetrathionate as a respiratory electron acceptor, a property that has been used empirically in clinical laboratories since 1923 to enrich for this pathogen in samples containing competing microbes [41]. However, tetrathionate respiration is of biological significance because it confers a luminal fitness advantage upon *S. enterica* in the inflamed intestine, thereby resulting in a disruption of the microbiota composition, which is characterized by an outgrowth of the pathogen [40]. The uncontrolled growth of *S. enterica* in the lumen of the inflamed gut enhances its transmission by the fecal oral route, thereby placing tetrathionate respiration under selection [42].

The finding that the inflammatory host response generates an exogenous electron acceptor in the gut lumen suggests that changes in the nutritional environment are a by-product of the antimicrobial activity of ROS and RNS. While ROS and RNS create a hostile environment in close proximity to the mucosal surface, these radicals are short lived and quickly react to form harmless oxidation by-products, such as tetrathionate. In addition to tetrathionate, a number of other oxidation by-products can drastically change bacterial growth conditions in the anaerobic environment of the large bowel. Nitric oxide (NO) generated by iNOS can react with superoxide radicals ($O_2^{\cdot-}$) produced by host NADPH oxidases to yield the bacteriocidal compound peroxynitrite ($ONOO^-$) [43,44]. Peroxynitrite is further converted to nitrate (NO_3^-) in a reaction catalyzed by carbon dioxide [45]. Through this mechanism, intestinal inflammation generates nitrate in the gut lumen [46,47] (Fig. 1B). Furthermore, ROS and RNS can oxidize organic sulfides, such as methionine, or tertiary amines, such as trimethylamine (TMA), to form the respective *S*-oxides and *N*-oxides [48,49]. Nitrate, tetrathionate, *S*-oxides and *N*-oxides are harmless oxidation products that can serve as exogenous electron acceptors for anaerobic respiration (reviewed in [50]). Thus, a by-product of releasing bacteriocidal ROS and RNS during inflammation is the generation of a cocktail of host-derived exogenous electron acceptors that enable microbes to perform anaerobic respiration.

The nitrate/nitrite redox couple has a high standard redox potential ($E^\circ = 433$ mV), which is second only to that of the oxygen/water redox couple ($E^\circ = 818$ mV) (reviewed in [51]). Under anaerobic conditions, nitrate is therefore the most potent electron acceptor for energy production (reviewed in [50]). Among the phylogenetic groupings that are present within gut-associated microbial communities, genes encoding nitrate reductase activity are found most commonly within genomes of the facultative anaerobic Enterobacteriaceae [52]. In contrast, genes encoding nitrate reductase activity are notably absent in genomes of obligate anaerobic bacteria belonging to the Bacteroidia and Clostridia [52]. The generation of host-derived nitrate during inflammation is thus predicted to favor growth of Enterobacteriaceae, because members of this family happen to be more likely to encode the enzymes required for nitrate respiration.

Direct evidence that inflammation-derived nitrate boosts growth of Enterobacteriaceae comes from studies on two of its representatives, *S. enterica* and *E. coli*. Colitis induced by *S. enterica* infection leads to the production of nitrate in the gut lumen, which in turn increases growth of the pathogen by nitrate respiration [53]. To enhance its growth in the gut lumen, *S. enterica* uses motility and chemotaxis to actively seek out metabolic niches that contain respiratory electron acceptors, such as nitrate and tetrathionate [54]. Similarly, nitrate generated by iNOS during chemically-induced or genetically-induced colitis markedly increases the luminal abundance of *E. coli* by supporting growth of this commensal microbe through nitrate respiration [47]. Host-derived nitrate is also generated during low-level intestinal inflammation induced by oral antibiotic treatment, thereby conferring a nitrate respiration-dependent fitness advantage upon commensal *E. coli* [55]. It has been proposed that the generation of nitrate in the intestinal lumen is one of the mechanisms by which antibiotic treatment reduces colonization resistance against commensal *E. coli* and other Enterobacteriaceae [56]. Luminal growth of *E. coli* is likely fueled further by respiration of other inflammation-derived electron acceptors, such as *S*-oxides and *N*-oxides [40]. In summary, the generation of exogenous electron acceptors by the inflammatory host response provides Enterobacteriaceae with a decisive luminal fitness advantage, which results in their uncontrolled expansion in the large bowel.

Why inflammation-derived electron acceptors favor Enterobacteriaceae

Exogenous electron acceptors enable Enterobacteriaceae to use strategies for maintaining redox balance, generating energy and acquiring carbon for biosynthesis of primary metabolites that are fundamentally different from those employed by Bacteroidia and Clostridia, thereby creating a new metabolic niche for these facultative anaerobic bacteria (Fig. 1). There are three advantages the metabolic strategy of Enterobacteriaceae has over that employed by Bacteroidia and Clostridia.

First, anaerobic respiration enables Enterobacteriaceae to balance their redox sheet by transferring electrons from NADH onto respiratory electron acceptors, such as nitrate, thereby preserving phosphoenolpyruvate for anabolic reactions. In contrast, fumarate respiration and fermentation performed by Bacteroidia and Clostridia are accompanied by the removal of metabolically valuable phosphoenolpyruvate to form metabolic end products to maintain redox balance.

Second, Enterobacteriaceae can use metabolic end products of Bacteroidia and Clostridia, such as formate or hydrogen, as electron donors to produce energy by anaerobic respiration (reviewed in [50]). This process can be performed solely for the purpose of energy production. For example, the transfer of electrons from hydrogen onto nitrate is independent of both carbon acquisition and maintaining redox balance. The finding that hydrogen enhances luminal growth of *S. enterica* in a mouse model [57], suggests that the ability to use metabolic end products of Bacteroidia and Clostridia for the sole purpose of producing energy provides a fitness advantage.

Third, anaerobic respiration broadens the spectrum of compounds that can serve as carbon sources in the anaerobic environment of the distal gut. During *S. enterica* infection,

intestinal contents are removed by the flushing action of diarrhea, thereby limiting microbial nutrition to compounds available in the mucus layer. Phosphatidylethanolamine, the most abundant phospholipid of enterocytes [58], is released into the mucus by sloughing and subsequently fermented by the microbiota, which produces ethanolamine as one of the end products (Fig. 1A). In the presence of an exogenous electron acceptor, such as tetrathionate, *S. enterica* can grow anaerobically on ethanolamine as the sole carbon source [59]. The presence of tetrathionate in the inflamed gut enables *S. enterica* to utilize ethanolamine to boost its luminal growth [60]. Collectively, these data suggest that the ability to utilize fermentation end products of Bacteroidia and Clostridia as a carbon source for the biosynthesis of proteins, carbohydrates, lipids and nucleotides confers a growth advantage upon *S. enterica*.

Conclusions

Intestinal inflammation has an impact on microbial metabolism through two different mechanisms. The first is related to the activity of antimicrobial proteins, which are released into the intestinal lumen during inflammation. The presence of these antimicrobial proteins favors growth of bacteria that are resistant against host nutrient withholding mechanisms [5,8]. As a result, nutritional immunity can alter the outcome of a competition between individual microbes [30]. A mechanistic understanding of these processes might facilitate the rational design of probiotics with increased capacity to exclude enteric pathogens.

The second mechanism influencing microbial metabolism is the generation of exogenous electron acceptors as a by-product of the inflammatory host response. In the anaerobic environment of the healthy gut, nutrient acquisition strategies of Enterobacteriaceae are inferior to the metabolic tactics employed by Bacteroidia and Clostridia (reviewed in [61]). However, the generation of exogenous electron acceptors by the inflammatory host response creates a new metabolic niche in the gut lumen. Unlike the obligate anaerobic Bacteroidia and Clostridia, the facultative anaerobic Enterobacteriaceae possess the enzymes to take advantage of the novel opportunities for generating energy, maintaining redox balance and acquiring carbon that become available in this new niche. As Enterobacteriaceae fill the metabolic niche created by intestinal inflammation, their relative abundance within the community increases and the resulting disruption of a balanced microbiota composition is known as dysbiosis. A dysbiosis characterized by a bloom of Enterobacteriaceae in the lower gastrointestinal tract is the ecological pattern observed most consistently in studies describing the changes in microbial communities that accompany gut inflammation (reviewed in [62]). A mechanistic understanding of the mechanisms responsible for dysbiosis identifies anaerobic respiration as a potential target for intervention strategies aimed at restoring a balanced community structure to improve health.

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Abbreviations

DUOX2	dual function NADPH oxidase 2
IFN-γ	gamma interferon
IL	interleukin
iNOS	inducible nitric oxide synthase
MPO	myeloperoxidase
NADH	nicotinamide adenine dinucleotide
NADPH	nicotinamide adenine dinucleotide phosphate
NOX1	NADPH oxidase 1
PHOX	phagocyte NADPH oxidase
RegIIIβ	regenerating islet-derived 3 beta
RNS	reactive nitrogen species
ROS	reactive oxygen species
SOD	superoxide dismutase
TCA	tricarboxylic acid
TMA	trimethylamine

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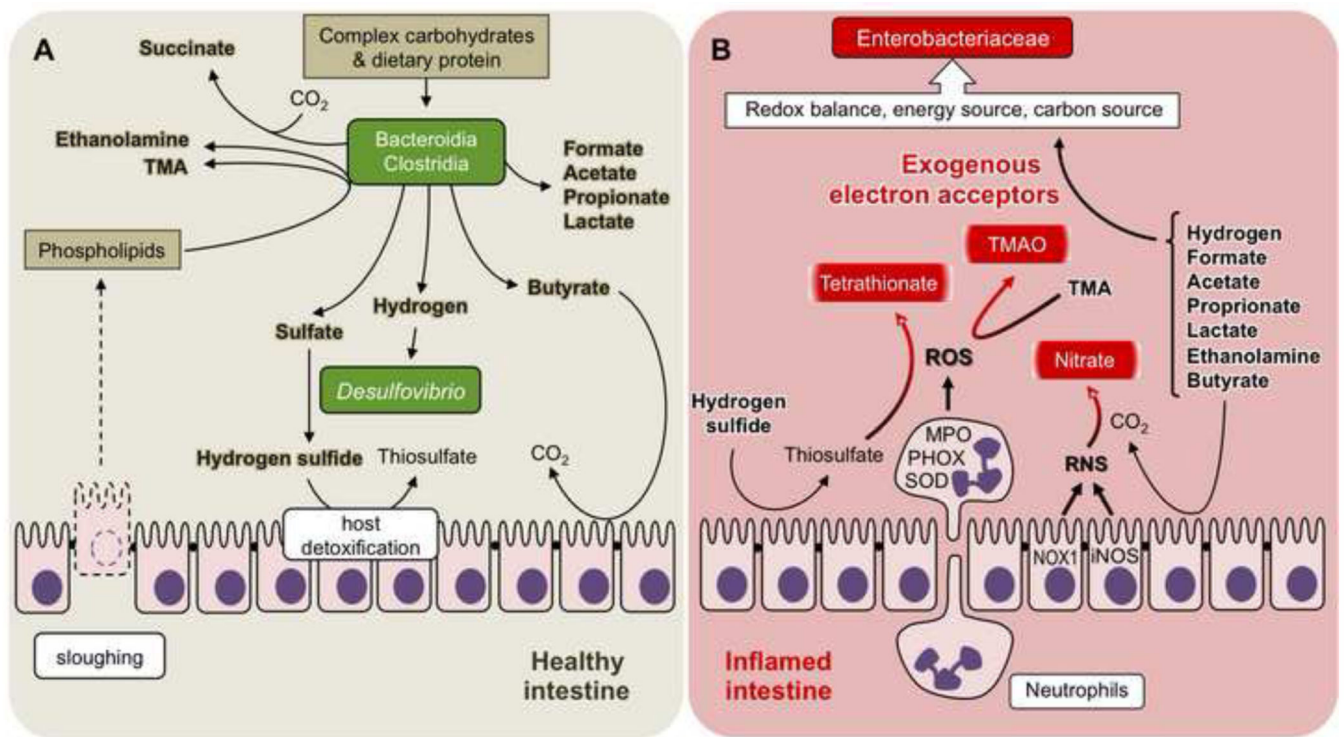


Figure 1.

The inflammatory host response creates a new metabolic niche in the intestine.

(A) The metabolic niche occupied by obligate anaerobic bacteria in the healthy intestine. Clostridia and Bacteroidia degrade complex carbohydrates and proteins in the distal gut to form a variety of metabolic end products, which accumulate in the lumen [32]. Furthermore, the head groups of phospholipids released by sloughing are degraded to TMA and ethanolamine [63]. Hydrogen produced by Clostridia and Bacteroidia fuels the growth of sulfate-reducing bacteria (*Desulfovibrio*), which produce hydrogen sulfide [35-37], a toxic gas oxidized by colonocytes to form harmless thiosulfate [38,39]. Butyrate is a fermentation product of obligate anaerobic bacteria that serves as nutrient for colonocytes [34]. (B) The metabolic niche occupied by facultative anaerobic Enterobacteriaceae in the healthy intestine. Reactive oxygen species (ROS) and reactive nitrogen species (RNS) generated by the inflammatory host response oxidize luminal compounds (TMA and thiosulfate) to form exogenous electron acceptors (trimethylamine *N*-oxide [TMAO] and tetrathionate, respectively) [40,48,49]. Some RNS species are converted into the exogenous electron acceptor nitrate in a reaction catalyzed by carbon dioxide (CO₂) [45]. The presence of exogenous electron acceptors enables Enterobacteriaceae to utilize microbiota-derived metabolic end products to generate energy, maintain redox balance and acquire carbon for the biosynthesis of primary metabolites.