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Review

The Role of the Intestinal Microbiome in Multiple Sclerosis—Lessons to Be Learned from Hippocrates

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Simple Summary: Multiple sclerosis is one of the long-term human diseases that significantly affects quality of life through the decline of muscle strength, problems in the coordination of muscle functions leading to disability or the decline of visual and cognitive functions. Although extensive research in the field of multiple sclerosis identified the abnormal function of the patients' immune systems as the primary mechanism underlying the disease, the ultimate cause of the autoimmune nature of multiple sclerosis, remained unknown. Recent research, however, shed light on the possible role of the bacteria living in the human gut, termed the human intestinal microbiome, in the development of multiple sclerosis. In this article, we review our current knowledge on the interplay between the intestinal microbiome and the immune mechanisms involved in the development of multiple sclerosis.



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Abstract: Based on recent advances in research of chronic inflammatory conditions, there is a growing body of evidence that suggests a close correlation between the microbiota of the gastrointestinal tract and the physiologic activity of the immune system. This raises the idea that disturbances of the GI ecosystem contribute to the unfolding of chronic diseases including neurodegenerative pathologies. Here, we overview our current understanding on the putative interaction between the gut microbiota and the immune system from the aspect of multiple sclerosis, one of the autoimmune conditions accompanied by severe chronic neuroinflammation that affects millions of people worldwide.

Keywords: intestinal microbiome; multiple sclerosis; macrophages; regulatory T lymphocytes

1. Introduction

Chronic inflammatory human diseases present a growing medical challenge to the healthcare systems in developed countries [1]. Despite the diversity of the involved tissues and organs, one of the possible underlying mechanisms is the disturbed control of the immune system that develops autoimmunity. Although the molecular details are still not clear, recent research suggests that the microbiota of the gastrointestinal (GI) tract plays an important role in the physiologic activity of the immune system, raising the idea that disturbances of the GI ecosystem contribute to the unfolding of autoimmune conditions. Here, we overview this putative interaction in the context of multiple sclerosis (MS), one of the autoimmune conditions accompanied by severe chronic neuroinflammation affecting millions of people worldwide.

2. Multiple Sclerosis

Multiple sclerosis (MS) is a common multifactorial chronic neurodegenerative disorder involving the central nervous system (CNS) affecting around 2 million people worldwide [2,3]. The primary symptoms of MS are directly related to the demyelination of axons, consisting sensory issues and the impairment of motor functions like ataxia and weakness [4]. Primary symptoms can lead to secondary complications including recurrent urinary tract infections due to impaired neuromuscular functions of the bladder or psychological problems like depression [5,6]. Based on the initial presenting symptoms, the disease courses are grouped into four major classes including the relapsing–remitting, primary progressive, secondary progressive and clinically isolated forms [7]. Currently, there is no cure available for MS, so mainstream therapeutic approaches focus on modifying disease progression by reducing the number of flareups and treating individual symptoms to improve patients' life quality [8].

The pathogenesis of MS is believed to be autoimmune, involving local inflammation and lymphocytic destruction of oligodendrocytes, the myelinating cells of the CNS (Figure 1) [9]. Myelin is a protective sheath that covers the neuronal axons and is involved in the accelerated conduction of the action potential [10]. The autoimmune destruction of myelin impairs communication between neurons causing sensory, motor and cognitive dysfunctions [11]. The death of oligodendrocytes in MS occurs via multiple pathways (Figure 1). One of these is binding of the Fas ligand to its receptor on the surfaces of oligodendrocytes, leading to their caspase-8-mediated apoptosis. Indeed, immunohistochemistry displayed an increased expression of Fas on oligodendrocytes and Fas ligand on microglia cells in MS [12]. Another mechanism of oligodendrocyte death is through TNF- α mediated necrosis [13]. In accordance, a correlation was noted between the TNF- α concentration in the cerebrospinal fluid (CSF) and the severity of the disease presentation in MS [14]. The destruction of the oligodendrocytes can also be mediated by IFN- γ activated caspase-11, as it was demonstrated in caspase-11 knockout mice, where cells remained resistant to cell death upon IFN- γ treatment [15]. The putative role of pro-inflammatory cytokines and the extrinsic apoptotic pathway in the destruction of the oligodendrocytes fuels the idea of the autoimmune nature of MS.

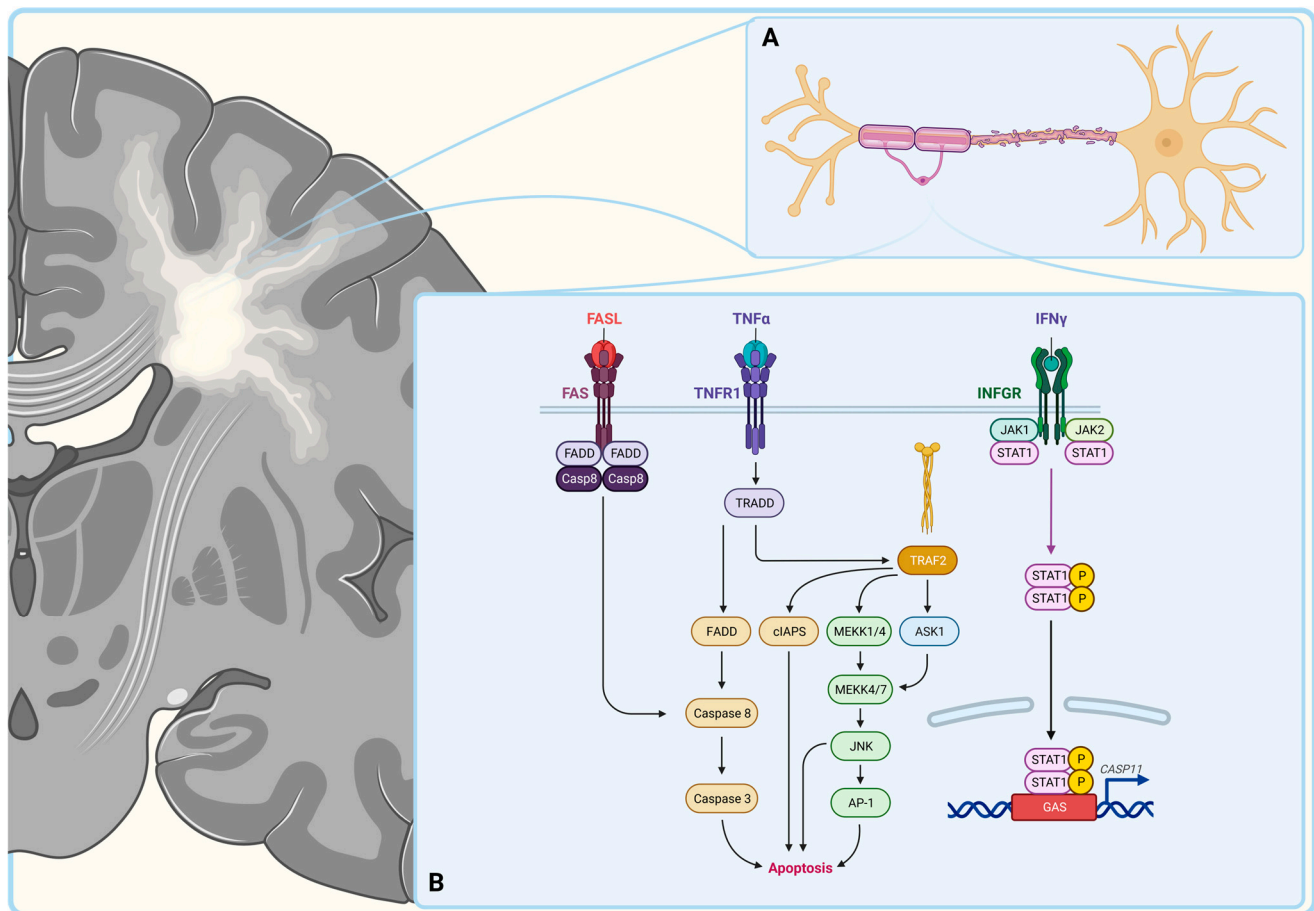


Figure 1. Pathological hallmark of multiple sclerosis. Multiple sclerosis is one of the demyelinating disorders with the leading pathological finding being the destruction of the myelin sheet around the axons (A). Oligodendrocytes can engage in apoptosis via the activation of multiple signaling pathways in response to the FAS ligand (FASL), Tumor necrosis factor alpha (TNF α) or Interferon gamma (IFN γ) (B). Created with [BioRender.com](https://www.biorender.com).

3. Autoimmunity in Multiple Sclerosis

Data indicate that upon the pathogenesis of MS, both innate and adaptive immune cells bypass the highly selective blood–brain barrier and become activated by myelin in a type IV hypersensitivity reaction [16]. It is believed that in MS patients, T lymphocytes are activated in the peripheral compartment and infiltrate the CNS leading to myelin and axonal damage [17]. The activation of autoreactive immune cells causes the secretion of cytokines, e.g., IL-1, IL-6 and TNF- α , compromising the blood–brain barrier (Figure 2) [18]. These pro-inflammatory cytokines upregulate the expression of adhesion molecules on vascular endothelial cells [19,20]. Lymphocytes bind to these adhesion molecules, migrate out of the vasculature and degrade the blood–brain barrier via matrix metalloproteases [21,22]. Early in the disease, macrophages dominate the infiltration followed by CD8⁺ and CD4⁺ T cells which, eventually, leads to general atrophy of the brain on imaging [23]. The majority of the CNS-infiltrating CD4⁺ T cells fall into the class of T_H1 and T_H17 cells, skewing the differentiation of CD4⁺ T cells away from the T_H2 phenotype [24]. As the disease progresses, B- and plasma cells begin to accumulate in the connective tissue of CNS as well forming ectopic follicles [25]. Although it is unclear if the presence of follicles causes the severe disease or if it is rather the consequence of disease progression, the association between follicle formation and disease progression has been noted, [26,27]. Accordingly, immune cells are found to be autoreactive against myelin basic protein and proteolipid protein, components of the myelin sheets, and immunohistochemistry analyses identified

C3d-, C9- and IgG-coated oligodendrocytes in the CNS of MS patients [28–30]. The role of autoreactive immune cells in the pathogenesis of the neuroinflammation in the CNS of MS patients has been supported by the findings that myelin basic protein-activated CD5⁺ T cells are able to provoke experimental allergic encephalomyelitis in healthy mice and that HLA-DR-positive macrophages are also present in the CSF of MS patients [31,32].

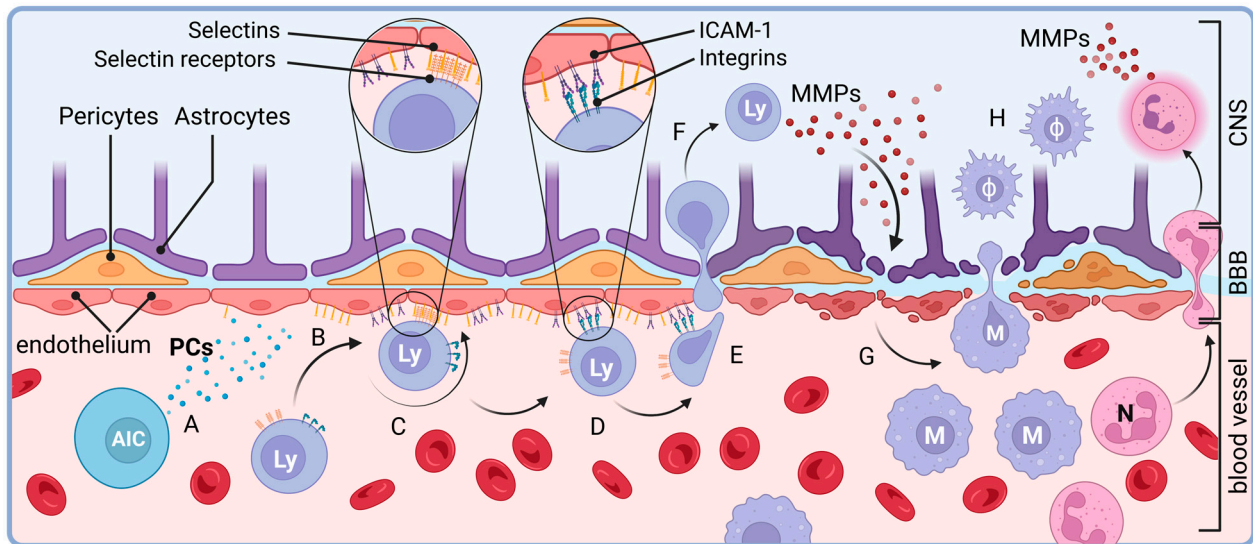


Figure 2. Pro-inflammatory macrophages invade the CNS in the initial phase of neuroinflammation in multiple sclerosis. Autoreactive immune cells (AIC)-released (A) pro-inflammatory cytokines (PC) induce the expression of adhesion molecules in endothelial cells that trigger recruitment (B) of lymphocytes (Ly), first, via weak Selectin (C) and, then, via firm, integrin-mediated (D) cell–cell interactions. The latter ones initiate diapedesis (E) of activated lymphocytes (F) into the interstitial space of CNS where secretion of matrix metalloproteinases (MMPs) damage *lamina basalis* of the blood–brain barrier (BBB). This leads to inflammatory injury of the cellular components of the blood–brain barrier including the endothelium, peri- and astrocytes alike. The damaged blood–brain barrier then recruits (G) monocytes (M) and neutrophil granulocytes (N). Following their immigration into the CNS, monocytes differentiate into pro-inflammatory macrophages (ϕ) and, with neutrophil granulocytes (N), initiate neuroinflammation (H). Created with [BioRender.com](https://www.biorender.com).

3.1. Macrophages

Besides monocytes recruited from the blood stream upon local inflammation, microglial cells, the tissue-resident macrophages of the CNS, are also believed to be key elements of MS pathogenesis. Tissue-resident macrophages are found locally in various tissues from birth where their tissue pools are maintained by in situ, macrophage colony-stimulating factor (M-CSF)/granulocyte macrophage colony-stimulating factor-dependent (GM-CSF) proliferation and play essential roles in immunosurveillance [33–35]. Embryologically, microglial cells are derived from primitive myeloid progenitors in the yolk sac, a distinct lineage compared to the hematopoietic stem cells, prior to the formation of the blood–brain barrier [36–38].

The putative role of microglial cells in the initial stages was established on the basis of the observation that the onset of the experimental autoimmune encephalomyelitis (EAE), the commonly used in vivo model of MS, was significantly delayed in a conditionally “paralyzed” microglial cell model [39]. It is believed that the activation of microglial cells leads to an oxidative burst causing free radical injury to the oligodendrocytes [40]. The progression from the initial stage, however, seems to be independent of microglial activation and rather dependent on the recruitment of monocytes from the blood [41]. This is mediated by the interaction between the monocyte chemoattractant protein-1 (MCP1) and the CC chemokine receptor 2 (CCR2) [42]. Indeed, macrophages are found in the CNS

of MS patients and EAE models but in the absence of CCR2 on monocytes, progression of the EAE fails [43–45].

Macrophages that are usually present in acute and active inflammation sites are characterized by the surface markers MRP14 and 27E10. Monocytes migrate from blood vessels to sites of inflammation, followed by their differentiation into macrophages or dendritic cells (reviewed in [46]). Macrophages, then, develop either a pro-inflammatory M1 phenotype or an anti-inflammatory M2 phenotype [47,48]. Although in chronic MS, macrophages are typically positive for surface markers characteristic for matured species, in a relapsing model of EAE, an increase in circulating GM-CSF-activated CD11b⁺, CD62L⁺ and Ly6C(hi)⁺ pro-inflammatory monocytes was observed prior to relapse, suggesting that proinflammatory cells invade the CNS from extra-CNS reservoirs upon relapses [44,49]. The M1 phenotype secretes pro-inflammatory cytokines such as IFN- γ , TNF- α and IL-6 as well as reactive oxygen species leading to the death of myelinating cells in the CNS [50]. Since the macrophage-mediated disease progression in MS is, at least in part, GM-CSF-dependent that produced by a subset of T helper lymphocytes, T cells are believed to play a central role in MS pathogenesis [51].

3.2. T Lymphocytes

T lymphocytes are derived from hematopoietic stem cells in the bone marrow that travel to the thymus for maturation [52,53]. This process gives rise to a naïve T cell pool that does not react to autoantigens but is ready to recognize their cognate foreign antigens to mediate immune reaction (reviewed in [54]). The central role of autoreactive T cells in MS emerged from findings that susceptible animals can be prompted to develop EAE through the active induction or adoptive transfer of autoreactive myelin-specific lymphocytes [55–57]. These antigen-specific lymphocytes undergo colony expansion in the regional node before migrating to the central nervous system (CNS) and forming inflammatory centers, which result in symptoms analogous to those observed in MS. The concept of the central role of T cells in MS is further underpinned by the International Multiple Sclerosis Genetics Consortium showing that the strongest association with genetic risk factors is with the HLA-DRB*1501 and HLA-A*0301 MHC alleles, key mediators of T cell antigen recognition [58].

The two, apparently, most prominent fractions of T lymphocytes in MS lesions are the CD4⁺ Th1 and CD4⁺ Th17 populations expressing IFN γ and IL-17, respectively [59,60]. The latter one is believed to not only maintain Th17 cells in an autocrine manner but also to promote pro-inflammatory reactions within the CNS (reviewed in [61]). Besides IL-17, Th17 cells also secrete IL-21 and -22 which are thought to contribute to the destruction of the blood–brain barrier, supporting the invasion of the CNS by activated immune cells [62].

Upon thymic education, a subtype of T cells, termed regulatory T cells (T_{reg}) and characterized by the CD4 and CD25 surface markers and the expression of the nuclear transcription factor FOXP3, also remain autoreactive, but that allows them to recognize and bind to autoantigens on the surface of other immune cells [63,64]. Indeed, T_{reg} cells help to maintain peripheral immunotolerance by inducing apoptosis in autoreactive T cells [65]. In the context of MS, depletion of T_{reg} cells suppresses the spontaneous recovery from EAE while an increase in T_{reg} cells is noted upon recovery from EAE [66]. In support of this concept, the induction of myelin oligodendrocyte glycoprotein-specific T_{reg} cells leads to the reduction of neuroinflammation in EAE models [67]. Data indicate that the recruitment of T_{reg} cells from the circulation upon remission in the EAE model is mediated via IL-4 [68]. The suppression of inflammation via T_{reg} cells is mediated via anti-inflammatory cytokines such as IL-10 [69]. IL-10 inhibits T cells by suppressing the co-stimulatory pathway for T cell activation via CD28 [70,71]. Indeed, EAE models with IL-10 knockout showed an increased release of pro-inflammatory cytokines [72]. In accordance, in MS patients, a functional defect in IL-10 secretion by T_{reg} cells was observed [73]. These observations fueled the agreement on the leading role of autoreactive T lymphocytes in MS.

3.3. B Cells

Although MS was considered as a primarily T cell dysregulation-mediated condition for years, recently this picture has been shifted toward a condition dictated by a disfunction between T and B lymphocytes. Indeed, the presence of oligoclonal immunoglobulins in the cerebrospinal fluid of MS patients and the positive effects of anti-CD20 antibodies in the clinical management of MS shed light on the importance of B cells in the pathogenesis of MS [74,75]. In secondary lymphoid organs, naïve B cells process antigens and load onto class II HLA complexes to present them for CD4⁺ Th lymphocytes, the key players in MS pathology [76]. Interestingly, the strongest genetic risk factor found in MS is the HLA-DRB*1501 allele that encodes for a class II MHC molecule [58]. In return, B cells respond to Th cell-secreted IL-21 which trigger their switch to IgG⁺ species [77]. It is noteworthy that this is the same cytokine that is secreted by Th17 cells and is believed to play a role in the destruction of the blood–brain barrier in MS [78]. Thus, one can speculate whether the HLA-DRB*1501 allele and/or autoreactive Th17 cells compromise the reciprocal interaction between B and T cells in MS patients. Indeed, data indicate that, besides the autoreactive T lymphocytes, MS patients harbor defects in peripheral B cell tolerance checkpoints, leaving polyreactive B cells behind in the circulation [79]. Although details of the underlying mechanisms are still to be elucidated, dysregulated B cells eventually begin to express high levels of CXCR3, enabling them to cross the blood–brain barrier in MS patients [80]. Data indicate that in the CNS, CXCR3⁺ species differentiate into antibody-secreting cells that support the recruitment of further CD4⁺ T cells to the CNS [81].

3.4. Neutrophil Granulocytes

It is noteworthy, however, that the CD4⁺ T cell-driven EAE models that established the T cell-centered view of MS pathogenesis neither reproduce the full-blown human disease nor necessarily display the potential role of other immune cell subtypes in MS. Indeed, IL-17, for instance, is one of the cytokines that promotes the migration of IL-1 β -producing neutrophil granulocytes to the CNS lymphatic system [82]. Although they were neglected initially in the context of this disease, neutrophil granulocytes are, now, believed to be potentially involved in the pathogenesis of MS as well. Indeed, neutrophil granulocyte products are detectable in the CSF during the onset of EAE, suggesting that the innate immune system also plays a role in the pathogenesis of MS [83]. This concept is further supported by findings that EAE animals show a delayed onset of EAE upon depletion of neutrophil granulocytes and that granulocyte macrophage-colony stimulating factor receptor knockout mice are resistant to EAE [84,85]. Moreover, the induction of genes encoding chemokines CXCL1 and CXCL2, which are known to be upregulated in the CNS during MS and are believed to orchestrate the chemotaxis of neutrophil granulocytes to the CSF and CNS in EAE, has been shown to be mediated by encephalitogenic CD4⁺ T_H17 cells, the core players of MS pathogenesis [86]. In accordance, genetic ablation of the major receptor for CXCL1 and CXCL2 (CXCR2) abrogates EAE in mice, but the transplantation of CXCR2⁺ polymorphonuclear leukocytes into CXCR2^{-/-} animals is sufficient to restore susceptibility to EAE [86]. In humans, markers of the primed state of neutrophil granulocyte activity, like their elevated number, reduced apoptotic activity, higher expression of TLR-2, fMLP receptor, IL-8 receptor and CD43, enhanced degranulation and increased levels of neutrophil extracellular traps in serum, have also been documented, depicting a complex autoreactive status of the CNS in MS and fueling intensive investigations for factors involved in the dysregulation of immune cells upon MS [87].

4. Microbiome as a Regulatory Factor of the Immune System

MS epidemiology suggests that these include ethno-genetic factors since MS is more prevalent in Caucasians compared to individuals of African or Asian descent [88]. The highest prevalence is observed in Northern Europe, particularly in regions like the British Isles and Scandinavia, as well as areas initially settled by migrants from these regions, including North America, Australia, and New Zealand [89]. These epidemiologic data

suggest a correlation between the prevalence of MS and the amount of winter sunlight, showing lower MS prevalence in regions with abundant sunlight [90]. This correlation, however, is not strong enough to consider sunlight exposure as the sole pathogenic factor of MS. Indeed, regions with very limited sunlight exposure, like the far north of Norway, show lower incident rates than that of Scotland or England and only half that of landlocked Oppland further south [91]. This peculiar pattern suggests the role of further ethno-genetic and/or geographic factors. Since latitude remains the most robustly associated factor of MS risk, inquiries into the role of photobiology linked vitamin D intake, particularly through multivitamin supplements, with a lower risk of MS [92–94]. A model that integrates vitamin D into genetic and environmental susceptibility to MS posits that vitamin D epigenetically modifies genes crucial in brain development, immune system function, axonal resilience, and immunological tolerance [90]. However, in a study assessing the effect of vitamin D supplementation on patients with relapsing–remitting multiple sclerosis (RR-MS), intriguing findings emerged, since vitamin D levels of RR-MS participants turned out to be very similar to that of the healthy participants both before and after 90 days of supplementation [95]. Instead, it was revealed that MS participants exhibited differential gut microbiota composition when compared to the healthy controls prior to supplementation of vitamin D. Specifically, they had lower levels of *Bacteroidaceae* and *Faecalibacterium*, and a higher abundance of *Ruminococcus*. It is particularly noteworthy that untreated MS subjects demonstrated a significant increase in the *Akkermansia*, *Faecalibacterium*, and *Coprococcus* genera following vitamin D supplementation [96]. These findings also raise the question of whether diet may influence MS incidence and progression, linking increased MS incidence to populations with high saturated fat intake and low vitamin D levels [90].

The potential involvement of dietary factors in MS pathogenesis naturally raises questions about the role of the microbiome (Figure 3). Indeed, emerging evidence suggests that the gastrointestinal microbiome plays an intricate role in shaping both innate and adaptive immunity, factors that are believed to be critical in the pathogenesis of MS. Studies on germ-free animals showed fundamental disturbances in the lymphoid tissue architecture and functions [97]. In the absence of the gut microbiota, IgA-mediated mucosal immunity is significantly reduced [98]. This is, at least in part, due to dendritic cells of the *lamina propria* that have been shown, sampling the mucosal layer of the intestinal lumen for microbial antigens both directly or through cellular elements of the gut epithelium like the Goblet cells, regulating immunoglobulin-producing B cells and T_{reg} lymphocytes alike [99–101]. *Helicobacter hepaticus* colonization in mice has also been shown to support T_{reg} responses through retinoic acid receptor-related orphan receptor γ t-positive (ROR γ t⁺) species, a novel class of antigen-presenting cells [102,103]. The mucosal antigen presenting cells interact with the common members of the gut microbiome as well. The most immunodominant zwitterionic bacterial polysaccharide, polysaccharide A, has been shown to be internalized and presented on MHCII to activate CD4⁺ T cells [104]. Thus, one can speculate that the composition of the gut microbiome determines the development of the mucosal immune system. Indeed, studies involving microbiome isolates from healthy subjects and patients with inflammatory bowel disease (IBD) have revealed diverse dendritic cell cytokine responses, driven by differential engagement of host Toll-like receptors (TLRs) [105]. These characteristic cytokine profiles include interleukin 6 (IL-6), tumor necrosis factor α (TNF α), IL-10, IL-23 and IL-1 β in response to *Bacteroidetes* and *Proteobacteria* in mouse myeloid cells in vitro [105]. Besides dendritic cells, segmented filamentous bacteria have also been shown to promote the development of antigen-specific Th17 cell responses underpinning the impact of microbiota on the regulation of the adaptive immune system [106,107].

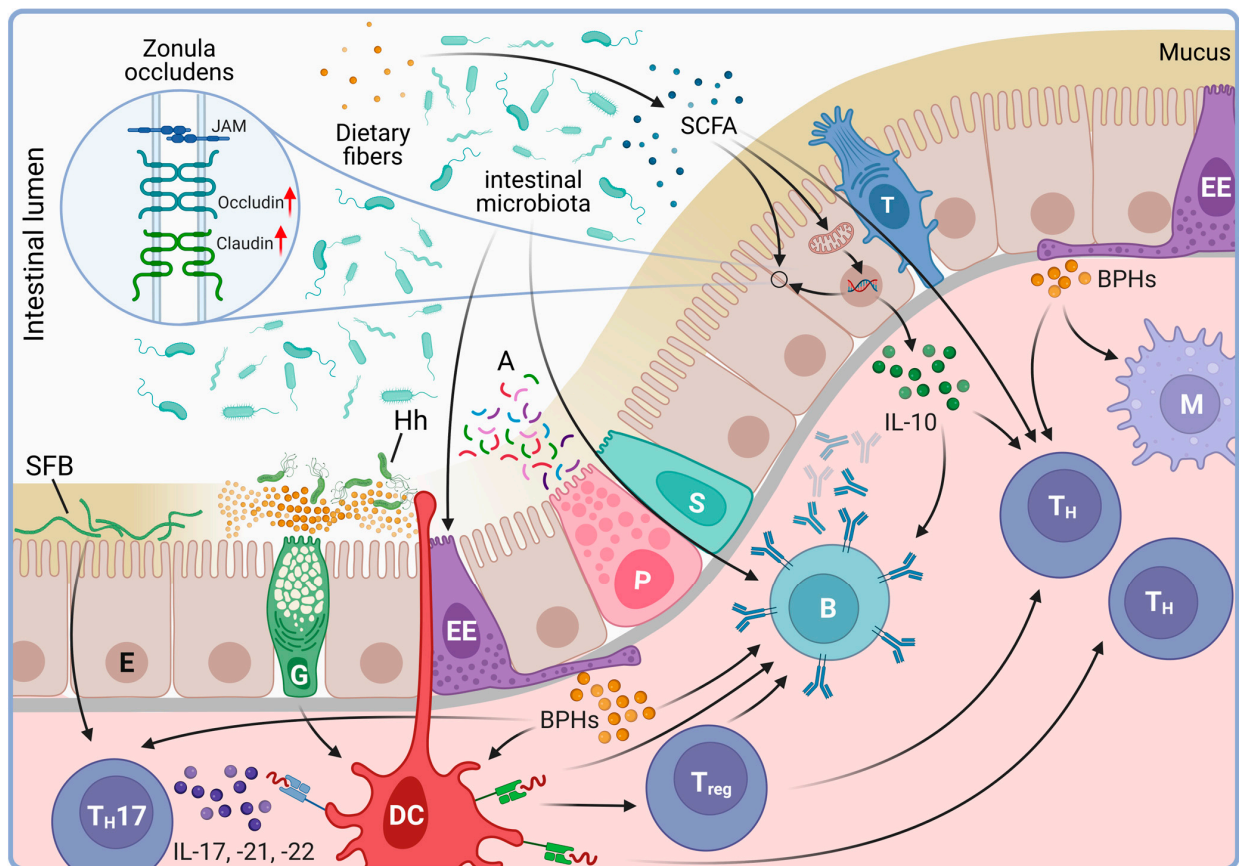


Figure 3. The gastrointestinal microbiome contributes to the functioning of the immune system. The gut microbiome and cellular elements of the intestinal epithelium have intricate interactions to maintain the barrier function of the gut lining. One of the critical components of this function is mediated by the tight junctions between enterocytes (E). Commensal germ metabolites, like short-chain fatty acids (SCFA), actively contribute to the upregulation of tight junction proteins. Parallel, enterocytes secrete cytokines that influence the immune cell population of the *lamina propria* including monocytes (M), antigen-producing cells (B) and helper (T_H) and regulatory (T_{reg}) T lymphocytes alike. Microbiota-derived antigens are also detected by enteroendocrine cells (EE), antigen presenting dendritic cells (DC) and enterocytes (E) by sampling the intestinal lumen directly. In return, enteroendocrine cells secrete bioactive peptide hormones (BPH), like serotonin, which, besides regulation of digestive functions of the enterocytes, affect B-, Th- and dendritic cell activities as well. The latter ones (DC), in contrast, can penetrate the intercellular space of the enterocyte lining and uptake microbial antigens directly from the mucus for presentation to immunocompetent cells of the epithelium. In the case of segmented filamentous bacteria (SFB), enterocytes can also mediate antigens to IL-17-producing T lymphocytes (T_H17) contributing to the regulation of the functionally critical Th17 pool of the intestines. For this effect, SFB need to contact enterocytes by penetrating the lower layers of the mucus. This physical barrier primarily consists of mucin produced by Goblet cells (G) which also sample the gut microbiota for antigens, e.g., the *Helicobacter hepaticus* (Hh), and delivers them to elements of innate immune cells of the *lamina propria* like the dendritic species (DC). Goblet cells are, at least in certain conditions, under the control of Tuft cells (T) that, in response to luminal antigens, for instance upon helminth invasion, produce IL-25. IL-25 activates type 2 innate lymphoid and T helper cells that, in response to IL-25, begin to produce IL-13. IL-13, in return, facilitates the commitment of epithelial stem cells (S) toward the production of Goblet- (G) and Tuft cells (T) [108]. The mucus also contains antimicrobial peptides (A) secreted by various elements of the epithelium including Paneth cells (P), which usually present in close vicinity of epithelial stem cells (S) and that, at least in part, are under the indirect control of dendritic cells via cytokines including IL-22 [109]. Created with [BioRender.com](https://www.biorender.com).

Microbiome-produced metabolites, like short chain fatty acids, vitamin D3 or the trimethylamine N-oxide (TMAO), also seem to have significant effects on the immune system [110–112]. Of these, the SCFA butyrate is believed to have the highest biological importance. Acting as an energy source for the intestinal epithelium, butyrate contributes to the maintenance of the physiologic hypoxia of the intestinal epithelium, stabilizing the transcription factor hypoxia inducible factor 1 (HIF1) that, in turn, upregulates genes that maintain integrity of the epithelial barrier [113]. Butyrate also functions as a histone deacetylase inhibitor, and as such, it is believed to exert anti-inflammatory properties by modulating pro-inflammatory cytokine production via the inhibition of nuclear factor κ B (NF- κ B) [114,115]. Other metabolites, like indole, attenuate the intestinal immune system by fortifying tight junctions in the epithelium via the upregulation of junction proteins like Claudin and direct induction of anti-inflammatory cytokines like IL-10 [116].

TMAO, in contrast, directly induces pro-inflammatory mediators like TNF α , NLRP3 inflammasome, mitochondrial reactive oxygen species (ROS), and NF- κ B, while reducing anti-inflammatory regulators such as IL-10 [117–119]. In accordance, TMAO triggers macrophage polarization toward the immunostimulatory M1 phenotype and enhances effector T cell functions, key factors of the pro-inflammatory state of the immune system [119]. These findings further support the idea of the potential relationship between the intestinal microbiota and the pathogenesis and progression of disorders associated with a dysregulation of the immune system like multiple sclerosis.

4.1. The Intestinal Microbiota

The first recognition of the microbial presence in the GI tract of living animals dated back to the mid-1800s [120]. The microbial taxa present in the special environment of the human GI tract is a complex ecosystem that presents a collection of microbes and their genomes referred as the gastrointestinal microbiome [121]. Its characterization began in the early 20th century leading to the isolation of *Escherichia coli* by Alfred Nissle and the ongoing work depicts an intricate ecosystem that includes hundreds of microbial species [122]. Indeed, microbial communities have been identified along the entire GI tract, although microbiome composition of distinct sections of the GI tract is differential [123] (Figure 4). In general, the vast majority (>99%) of the identified species belong to five bacterial phyla including the *Firmicutes*, *Actinobacteria*, *Bacteroidetes*, *Proteobacteria* and *Fusobacteria* (Figure 4) [124,125]. The microbiota of the throat, that mostly constituted of the *Streptococcus*, *Prevotella*, *Actinomyces*, *Gemella*, *Rothia*, *Granulicatella*, *Haemophilus*, and *Veillonella* genera, shows the lowest phylotype diversity and inter-individual variety compared to the microbiota of the stomach or the lower GI tract [126]. Indeed, *Streptococcus*, *Actinomyces*, *Prevotella* and *Gemella* are the most abundant in the stomach [127]. In contrast, in fecal samples, representing the lower GI tract, genomic analyses revealed a minimum of 109 species from 8 phyla, 18 families, 23 classes, 38 orders and 59 genera [128]. According to current data, in fecal samples of healthy individuals, the phylum *Firmicutes* dominates (approx. 40%) [129]. The majority of the *Firmicutes* belong to the *Clostridia* class with frequent representation of the *Clostridium*, *Eubacterium* and *Ruminococcus* genera. The latter includes the most abundant species, the *Ruminococcus bicirculans*, that alone makes up approximately 2.5% of all species in the lower GI tract. The second most dominant phyla, both with an approximate 20% share in fecal samples, are the *Actinobacter*, that most abundantly comprises the species *Bifidobacterium longum* from the genus *Bifidobacterium*, and the *Bacteroidetes* mostly represented by the species *Bacteroides fragilis* from the class of *Bacteroidia*. At the species level, *Escherichia coli* (1.87% of total species) and the *Enterococcus faecium* (0.04% of total species) of the *Enterobacterales* and *Lactobacillales* orders, respectively, are the overrepresented in the lower GI tract [129].

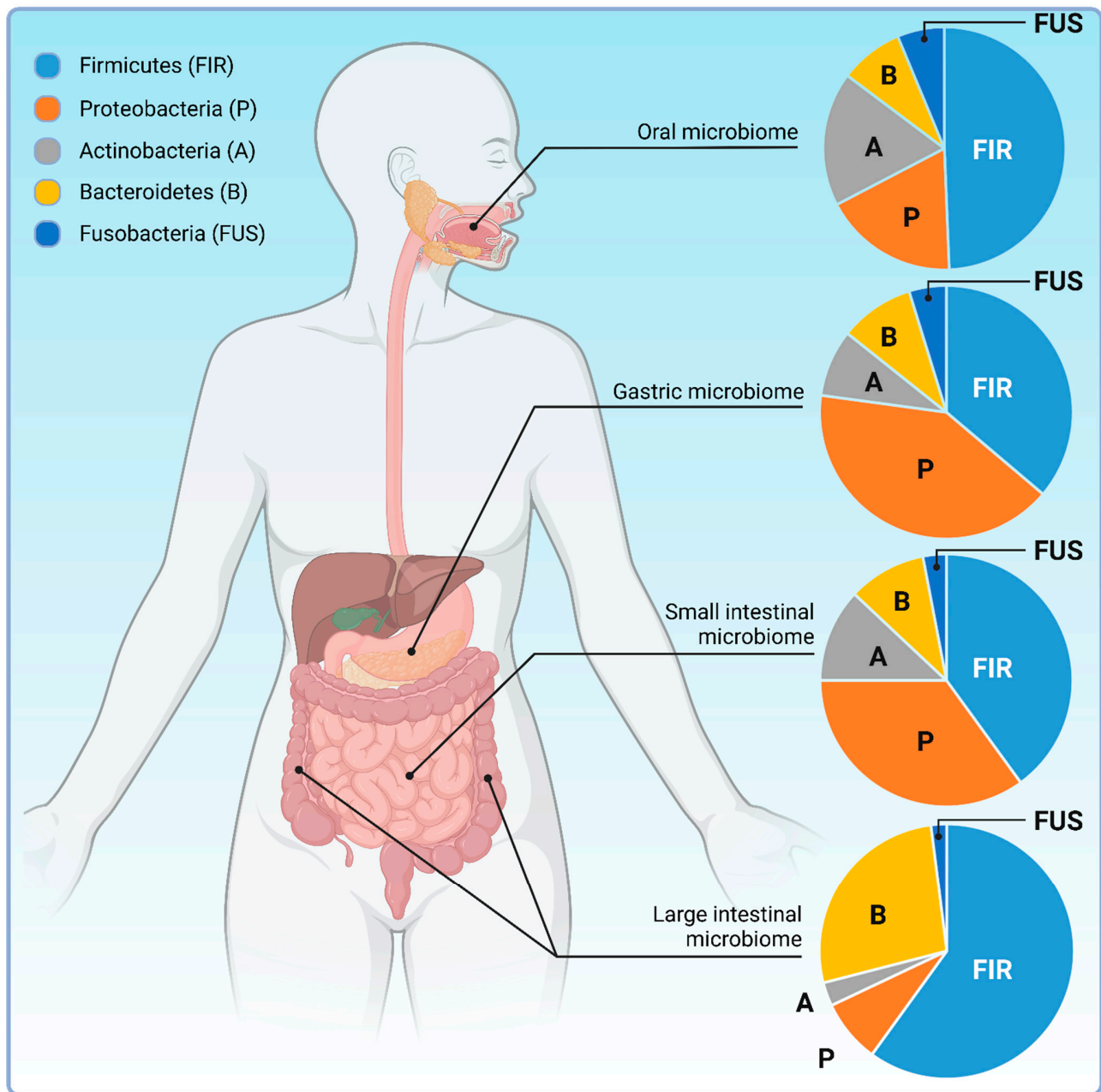


Figure 4. Spatially diverse composition of the gastrointestinal microbiome along the GI tract. Although predominant phyla are the same, proportional changes in microbiome diversity have been observed along the gastrointestinal tract of which the most characteristic is the reciprocal alteration in the presence of *Bacteroidetes/Firmicutes* and *Actino-/Proteobacteria* phyla in a cranio-caudal manner [129]. Created with [BioRender.com](https://www.biorender.com).

4.2. Origin of the GI Microbiome

While distinct sections of the GI tract show differential microbiome composition, spatial distribution of the identified species is not the only diversity found in the GI tract microbiome. Genetic analyses revealed that the GI tract microbiome also shows significant age-related variability and data indicate that gestational age heavily affects human GI microbiota composition [130]. Fecal samples of very low birth-weight infants showed that, between 25 and 30 weeks of postmenstrual age, the lower GI microbiome is primarily dominated by *Staphylococci*. This population, then, is overtaken by a *Bifidobacterium*-dominated microbiota from postmenstrual week 30 onwards, accompanied by a peak in the abundance

of *Enterococci* between postmenstrual weeks 30–35 and 45–50. In the case of term-born infants, the mouth flora was predominantly colonized by *Streptococci viridans* and *S. salivarius* by post-natal day 6 while the fecal flora showed the dominance of species from the classes of *Clostridia* and *Gammaproteobacteria* and the two most abundant species found were *Enterococcus faecalis* and *Escherichia coli* [130,131]. At the phyla level, it seems that delivery does not affect the colonization of the lower GI tract, since one month post-natal, fecal microbiota was found to be dominated by *Actinobacteria* independently of whether children were born via caesarean section (CS) or vaginal delivery (VD) [132]. Apparently, this changes at the later stages of the infant life since children's fecal samples are dominated by *Bacteroidetes* and several genera of the *Clostridia* class, for example the *Ruminococci*, by the end of the first two years of age. At lower taxonomy levels, however, some degree of differential microbiome composition was observed much earlier post-natal. The *Enterococcus* genus, for instance, was found in significantly higher relative abundance in CS infants compared to their VD counterparts. In contrast, VD infants showed a significantly higher level of *Bacteroidetes* during the first 12 months post-natal. Parallel, CS delivery was found to be associated with a significantly lower overall microbiota diversity, even though the mothers of the CS and VD children did not show similar differences in their microbiota diversity. It is noteworthy, however, that the composition of the infant gut microbiota is affected by the feeding regime as well. Indeed, *Bacteroidetes* and *Clostridia* are more prevalent in fecal samples of weaned children than that of the breast-fed infants, highlighting the complexity of the determining factors of the gut microbiome [133,134].

4.3. Dietary Links

Similar to infants, dietary aspects affect the composition of the gut microbiome in adults as well. Animal studies revealed that the intake of non-digestible complex carbohydrates of plant origin, also known as dietary fibers or microbiome-accessible carbohydrates, is one of the determining factors of the gut microbiota diversity [135]. In humanized rodent models, a low microbiome-accessible carbohydrates diet primarily reduces the abundance of *Bacteroidetes*, the second most abundant phylum of the healthy human gut microbiome. The consumption of nutrients, e.g., in broccoli, in contrast, increases the abundance of *Bacteroidetes* [136]. The effect of nutrients on the diversity of the gut microbiome, however, is not uniform. Indeed, while walnut consumption increases the relative abundance of genera *Faecalibacterium*, *Clostridium*, *Roseburia*, and *Dialister* of the *Firmicutes* phylum, it decreases the abundance of *Bifidobacteria* of the phylum *Actinobacteria* [137]. An interesting aspect of the dietary effects on the gut microbiome is that, according to certain experimental findings, they can lead to the complete eradication of some species of the gut microbiota. A dietary supplement of microbiome-accessible carbohydrates to low level microbiome-accessible carbohydrates-fed animals alone did not reverse gut microbiome diversity if not combined with fecal microbiota transplants (FMT) [67]. Considering the significant role of nutrients in its composition, it is not surprising that the gut microbiota also shows seasonal changes. A long-term dietary survey analyzing the microbiota composition of human fecal samples showed that members of the phyla *Firmicutes* and *Actinobacteria* are more abundant in the samples collected over wintertime while species from the phylum *Bacteroidetes* are more abundant in samples taken in the summer months [70]. This seems to be in accordance with the findings that the abundance of *Bifidobacterium longum*, the most abundant species of *Bifidobacteriaceae*, positively correlates with the vegetable, protein and soluble fiber intake, or that *Akkermansia*, members of the phylum *Verrucomicrobia*, were shown to be positively associated with saturated fat intake and negatively correlated with the amount of total polyunsaturated fatty acids present in the diet [128]. Data also indicate that overall calorie intake is another important determinant of the gut microbiota diversity, showing a positive correlation between the body weight, the waist circumference and the abundance of certain species of the gut microbiota like *Bacteriodes ovatus* [138]. Indeed, fat-restricted or carbohydrate-restricted diets were found to increase the relative abundance of *Firmicutes*

while decreasing the abundance of *Bacteroidetes*, underlining the importance of the dietary factors in the composition of the human gut microbiome [139].

4.4. Microbiome and Disease

Besides dietary factors, data suggest that the host's genetic background also determines the gut microbiota composition. Indeed, individuals carrying the HLA-DQ haplotype, a genetic condition that makes the individuals prone to Type I *Diabetes mellitus* (T1D), have a differential development course of their gut microbiome [140]. Indeed, the gut microbiome is dominated by members of the *Bacteroidetes* phylum in newborns during the first year post-natal, while, in children born with the HLA-DQ haplotype, the colonization of the gut by *Bacteroidetes* is delayed. In contrast, species representing the *Firmicutes* phylum peak early in the post-natal life of T1D-prone children, while they increase in abundance much later in non-HLA-DQ children. This early peak of colonization by *Firmicutes*, then, is accompanied by a depletion parallel with the increase in *Bacteroidetes* abundance, although, eventually, an overall reduction in the diversity of the gut microbiome of T1D-prone children by the age of year two was reported [140]. These data suggest that immune-related haplotypes interplay with the dynamics of the initial gut colonization, although it is not clear whether it is the gradually developing auto-immune environment that favors *Bacteroidetes* or the HLA-DQ haplotypes is the one that biases the gut microbiota toward *Bacteroidetes* which eventually induces the auto-immune condition leading to T1D.

The imbalance of *Bacteroidetes* and *Firmicutes* in the gut microbiome was documented in additional autoimmune pathologies too. In ulcerative colitis and Crohn's disease patients, for instance, depletion of species from the *Firmicutes* phylum was reported to be more dominant over the depletion of *Bacteroidetes* [141]. In systemic lupus erythematosus, the prototype of the autoimmune diseases, similar dysbiosis was observed, indicating that the interplay of the gut microbiota and the immune homeostasis of the host environment is not restricted to the gastrointestinal tract [142]. Indeed, depletion of the *Firmicutes* phylum and increased abundance of species of the *Bacteroidetes* was also reported in patients suffering from rheumatoid arthritis [143]. In addition, imbalanced gut microbiota composition was reported in patients suffering from the major depressive disorder (MDD) [144]. Moreover, the depressive condition seemed to be transferrable via the gut microbiome, indicating the pathogenic importance of the disturbed gut microbiome in neuropsychological disorders. These observations further support the idea that the complex effects of the gastrointestinal microbiome affect tissues that were traditionally considered to be functionally independent of the GI tract.

Indeed, there is a growing body of evidence suggesting that an alteration in the GI tract microbiota diversity correlates with chronic human pathologies of neuronal tissue as well. In relation to Parkinson's disease, for instance, the overall microbiota composition was found to be shifted with a depletion of butyrate-producing bacteria, and an overabundance of pro-inflammatory species like the *Collinsella*, *Desulfovibrio*, and *Oscillospiraceae* [145,146]. Even more interestingly, these alterations of the gut microbiome seem to be present in the long prodromal phase of Parkinson's disease, suggesting a causality between the two processes [145]. In addition, a similar trend was found in REM sleep behavior disorder [145]. The predicted functional profile showed an overall increase in fatty acids' fermentation to lactate and ethanol, and lower levels of deazapurine biosynthesis in both pathologies [145]. Reduced gut microbiome diversity has been reported in relation to Alzheimer's disease as well. A recent meta-analysis of related studies showed an overall decrease in species richness in the Alzheimer's disease gut microbiome, fueling the idea of the possible role of the GI tract microbiome in the pathogenesis of further neurodegenerative disorders [147].

5. The Intestinal Microbiome and Multiple Sclerosis

The regulatory role of the intestinal microbiome in shaping the immune system sheds light on its putative involvement in the onset and progression of multiple sclerosis. Accordingly, it turned out that the gut microbiome of MS patients showed dysbiosis mainly

affecting genera of both *Firmicutes* and *Bacteroidetes* (Tables 1 and 2) [148]. Interestingly, however, the trend of the observed changes seems to be independent of the phylogenetic classification. While *Bacteroidetes* genera *Pedobacteria* and *Flavobacterium* showed a higher abundance in patients with MS, other *Bacteroidetes* genera like the *Parabacteroides*, *Bacteroides*, and *Prevotella* were rather depleted. Moreover, it also seems that the diversity of the gut microbiome correlates with the disease course. Indeed, the small intestinal microbiota of patients suffering from RR-MS have an increase in the phylum *Firmicutes* and a decrease in the phylum *Bacteroidetes* during relapse compared to both healthy individuals and MS patients in remission [148–150].

Table 1. Genera of gut microbiota with increased abundance in multiple sclerosis.

Phylum	Genus
<i>Bacteroidetes</i>	<i>Pedobacteria</i>
	<i>Flavobacterium</i>
	<i>Dorea</i>
<i>Firmicutes</i>	<i>Balutia</i>
	<i>Streptococcus</i>
	<i>Mycoplana</i>
<i>Proteobacteria</i>	<i>Acinetobacter</i>
	<i>Pseudomonas</i>
<i>Actinobacteria</i>	<i>Eggerthella</i>
<i>Verrucomicrobia</i>	<i>Akkermansia</i>

Table 2. Genera of gut microbiota with decreased abundance in multiple sclerosis.

Phylum	Genus
<i>Bacteroidetes</i>	<i>Bacteroides</i>
	<i>Prevotella</i>
	<i>Parabacteroides</i>
	<i>Coprobacillus</i>
<i>Firmicutes</i>	<i>Lactobacillus</i>
	<i>Clostridium</i>
	<i>Anaerostipes</i>
	<i>Faecalibacterium</i>
<i>Proteobacteria</i>	<i>Haemophylus</i>
	<i>Sutterella</i>
<i>Actinobacteria</i>	<i>Adlercreutzia</i>
	<i>Collinsella</i>

Although results on the gut microbiome diversity of MS patients are not entirely consistent, altering these ratios in animal models has been shown to influence disease phenotype. Susceptible animals develop EAE through the active induction or adoptive transfer of autoreactive myelin-specific lymphocytes [151]. These antigen-specific lymphocytes undergo colony expansion in the regional node before migrating to the CNS and forming inflammatory centers, which results in symptoms analogous to those observed in MS. The relapsing–remitting model of spontaneously developing experimental autoimmune EAE has been used to demonstrate the involvement of microbiota in disease development [152]. In mice, the transfer of intestinal microbiota from individuals with MS has been linked to the spontaneous development of EAE (Figure 5). Additionally, when mice are colonized with the gut microbes of MS patients, the resulting experimental disease is much more severe [153,154]. Interestingly, human alterations in gut microbiome in MS include a decrease in *Prevotella* and an increase in *Streptococcus mitis* (*S. mitis*) and *S. oralis*, all believed to have roles in shaping the immune system. Indeed, the *Prevotella* reduction is accompanied by the expansion of T_H17 cells, while strains of the *S. mitis* group is known to positively affect the differentiation of T_H17 cells, key players in cell-mediated tissue damage and

autoimmunity in RR-MS patients [155,156]. Although the mechanistic relationship between these bacteria and the expansion of pro-inflammatory cell types like the T_H17 cells are not fully understood, one may speculate that their immunological effects are mediated by the metabolites they produce.

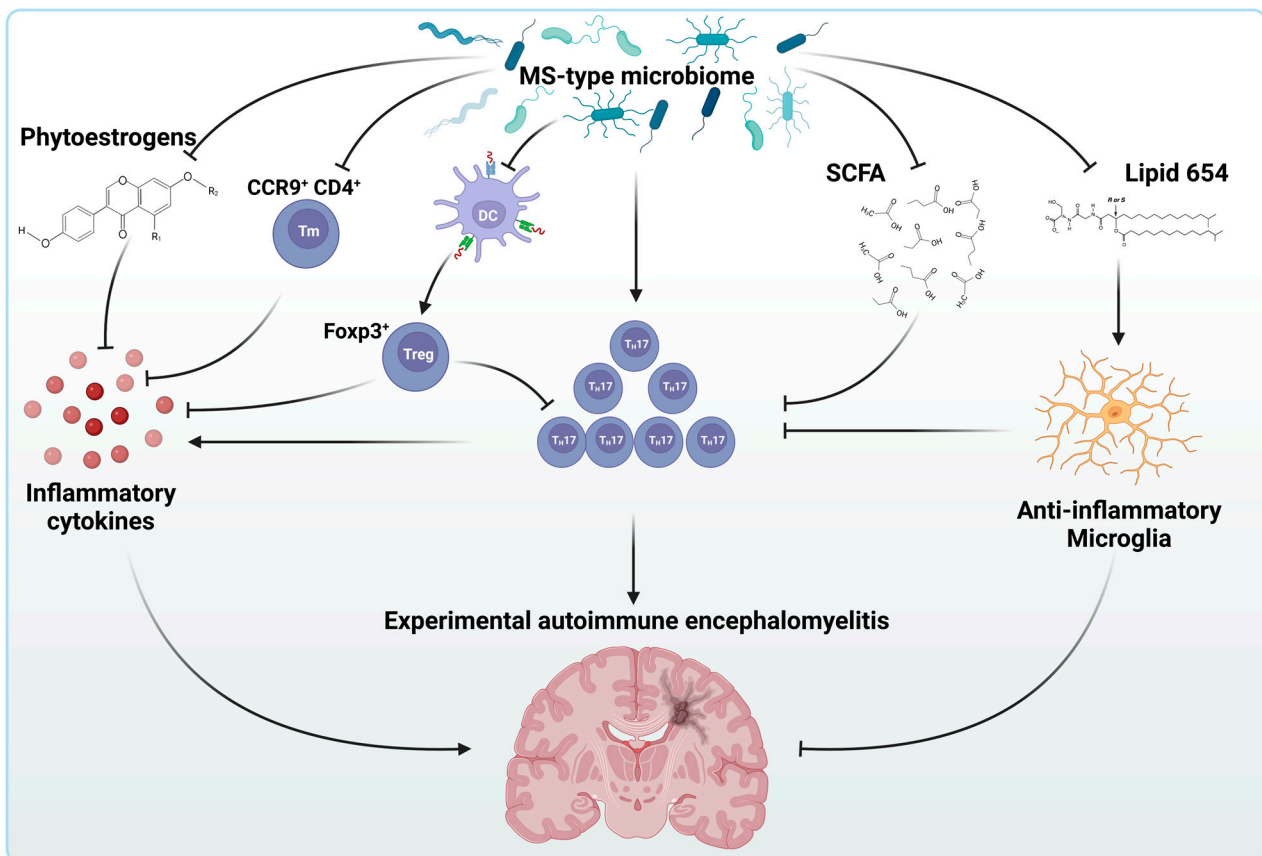


Figure 5. Dysbiosis of the gastrointestinal microbiome facilitates the proinflammatory state of multiple sclerosis. Alterations in the composition of the gastrointestinal microbiome influence the immunological status in multiple mechanisms in multiple sclerosis (MS). On one hand, MS-type microbiome directly facilitates differentiation and the expansion of T_H17 lymphocytes (T_H17) that play a central role in the maintenance of the inflammatory state of MS patients. Parallel, MS-type microbiome directly blocks the activity of anti-inflammatory species like the $CCR9^+$ memory T cells (Tm) or tissue-resident dendritic cells (DC). Blockade of the latter results in the failure of differentiation of $Foxp3^+$ regulatory T cells (T_{reg}) which results in the accumulation of pro-inflammatory cytokines and cellular species alike. On the other hand, the MS-type intestinal microbiome exerts pro-inflammatory effects via various microbial metabolites by blocking the physiologic metabolisms of compounds like the phytoestrogens, short-chain fatty acids (SCFA) or microbial lipids (Lipid 654). Created with [BioRender.com](https://www.biorender.com).

Indeed, it is widely believed that the anti-inflammatory effects of bacteria like *Faecalibacterium* or *Coprococcus* are primarily mediated by their butyrate production [157–159]. Similarly, *Prevotella* is a known propionate producer, that, along with acetate and butyrate, is a characteristic compound of the gut microbiome-produced short-chain fatty acids (SCFA) and is believed to be involved in the regulation of T_{reg} cells in peripheral compartments and the increase in the anti-inflammatory cytokine IL-10 [160–163]. To support this concept, dysbiosis of the gut microbiome in RR-MS patients has been observed to include the reduction of the *Clostridia* cluster XIV and IV accompanied by the reduction in the short-chain fatty acid production, also suggesting a widespread involvement of gut microbiota species

in the unfolding of the pro-inflammatory state of MS via, at least in part, the reduction in short-chain fatty acid production [149].

Lipid 654, another bacterial metabolite typically produced by *Bacteroidetes* species that acts as a Toll-like receptor-2 ligand, was also found to be significantly lower in MS patients [164,165]. According to recent data, lipid 654 alters the microglial response to pro-inflammatory cytokines like IFN β via shifting the microglial polarization to the rather anti-inflammatory M2 activation state [166]. Thus, the depletion of *Bacteroidetes* species in the MS gut microbiome might contribute to neuroinflammation via a reduction in lipid 654 production as well.

Besides the extensively studied *Firmicutes* and *Bacteroidetes* phyla, *Adlercreutzia* of the *Actinobacter* phylum has also been reported to be disturbed in MS patients' microbiomes [148]. *Adlercreutzia* is known to play a role in influencing anti-inflammatory responses by way of its connection to phytoestrogen metabolism. Phytoestrogens are plant-derived molecules that possess a chemical structure and biological activity similar to estrogen. Legumes (especially soybeans), fruits, whole grains, and other vegetables are the primary sources of these compounds. Bacteria, such as *Adlercreutzia*, through β -glucosidase, are responsible for converting phytoestrogens into monomers. In patients with RR-MS, a reduction in *Adlercreutzia* leads to a decrease in the conversion capacity of phytoestrogens. Consequently, the decrease in this bacterium results in an increase in oxidative stress and inflammatory cytokines, such as the monocyte chemo-attractant protein-1 (MCP-1/CCL2) and IL-6, which are normally elevated in MS [167].

The compromised MS gut microbiome has been reported to have an impact on another chemokine–receptor interaction between the T-cell C-C chemokine receptor type 9 (CCR9) and its ligand CCL25 [168]. This interaction is important for CD4⁺ memory T cell homing to the gut and, thus, the immunity of the small intestinal epithelium [169]. CCR9⁺ CD4⁺ T cells belong to the intraepithelial cohort of T cells that has been found to be protective against EAE [170]. Disruption of the CCR9-CCL25 interaction by the MS gut microbiota results in a depleted population of CCR9⁺ CD4⁺ T cells, suggesting another potential mechanism involved in the exacerbation of neuroinflammation in SP-MS patients.

The putative link between the immunologic mechanisms and the gut microbiota in MS has been indirectly supported by findings in MS patients treated with glatiramer acetate (GA) [95]. Although GA was originally developed to provoke experimental autoimmune encephalitis as an animal model of human MS, it turned out to be acting as an immunomodulator and has been approved by the FDA as a first line drug to treat relapsing MS [171]. GA is believed to reduce the secretion of pro-inflammatory cytokines (IL-2, IL-12, IFN γ , TNF) released by T_H1 cells, activate T_H2 suppressor cells to express anti-inflammatory cytokines (IL-4, IL-5, IL-13, IL-10, TGF- β) in the CNS and increase the enrichments of T_{regs} [172,173]. GA, however, has also been shown to modulate the composition of the gut microbiota in MS patients elevating the proportion of *Janthinobacterium* while decreasing the presence of *Eubacterium* and *Ruminococcus*, suggesting the a putative bidirectional interplay between the gut microbiota and the immune mechanisms involved in MS [95].

These observations inevitably raise the question of the causality in the interplay between the intestinal microbiome and the immune system in MS. The two main hypotheses of the pathogenesis of MS allows differential interpretation of the role of the intestinal microbiome in the development of the disease. The inside-out concept posits that MS is initiated by a primary lesion in the CNS, potentially triggered by factors like infection or primary neurodegeneration. This event leads to the release of self-antigens, provoking a response from autoreactive T and/or B cells. In this concept, one could speculate that the differential composition of the gut microbiome observed in MS patients is a consequence of a preexisting pro-inflammatory state of the immune system.

Conversely, the extrinsic or outside-in concept suggests that peripheral stimuli like microbial antigens resembling CNS molecular signatures, bystander activation of immunocompetent cells, novel autoantigen presentation, or recognition of secluded CNS antigens activate escaped autoreactive T cells, eventually leading to a neuroinflammatory state [174].

In this concept, disturbances in the gut microbiome might play the role of the initial antigen stimulus. This idea is, apparently, strongly supported by animal studies in EAE models. Indeed, the administration of differential combinations of antibiotics inhibits the development of EAE in susceptible animals. This disruption is linked to an increase in Foxp3⁺ regulatory T cells in the mesenteric and cervical lymph nodes which appear to be influenced by an elevated presence of the co-localized CD11c^{high}CD103⁺ dendritic cells. These specific DCs promote the transformation of naïve CD4⁺ T cells into Foxp3⁺ T_{reg} cells [175]. Parallel, an increased presence of IL-10-producing CD5⁺ B cells has also been observed in cervical lymph nodes. When splenic CD5⁺ B cells from antibiotic-treated mice were transferred to naïve recipient mice, and the recipients were immunized with MOG35–55 one day post-transplantation, a significant reduction in EAE disease score was noted. In addition, this reduction seems to be associated with a shift from a Th1/Th17 cytokine profile to a Th2 cytokine profile [176]. To further support the role of the gut microbiome in the extrinsic model of MS, the transplantation of the intestinal microbiota of MS patients into susceptible mice has been observed to be associated with the spontaneous onset of EAE. Moreover, when mice are colonized with microbes sourced from the gut of MS patients, the resultant experimental disease manifests with increased severity, suggesting a causative role of the microbiome in the development of MS in susceptible individuals [153,154].

Considering its complexity, one can speculate that the intestinal microbiome can provide a generous source of non-self biocompounds that could serve by triggering antigens for autoreactive species. Research identifying the guanosine diphosphate-L-fucose synthase as an autoantigen for CD4⁺ autoreactive T cells supports this idea [177]. This enzyme is involved in the nucleotide–sugar biosynthesis and catalyzes the conversion of GDP-4-dehydro-6-deoxy-D-mannose to GDP-fucose which serves as substrate for fucosyltransferases to the mediated fucosylation of oligosaccharides, glycoproteins, and glycolipids [178]. Although, in mammals, fucosylation plays a critically important role in a number of physiologic processes, in prokaryotes fucosylation is less common. Still, it has been demonstrated that certain bacteria of the gastrointestinal tract, like oral commensals that are associated with infected endocarditis, such as *Porphyromonas gingivalis*, *Actinobacillus actinomycetemcomitans* and *Eikenella corrodens* or the gastric *Helicobacter pylori*, exploit fucosylated structures on bacterial lipopolysaccharides (LPS) [179]. Indeed, in the latter, for instance, molecular mimicry through fucosylated LPS enhances *H. pylori* adhesion to the gastric epithelium and modulates hosts' immune responses to aid adaptation effectively to its niche and maintain a persistent infection [180]. Interestingly, the fucosylated polysaccharide chains of *H. pylori* LPS induce the production of auto-antibodies, causing antigastric autoreactivity and, consequently, tissue damage in the gastric mucosa [181].

Thus, one can speculate that the CD4⁺ T lymphocyte-mediated immune reaction against the guanosine diphosphate-L-fucose synthase is, primarily, a protective measure against potentially pathogenic invading microbes in the gastrointestinal tract that, eventually, turns into autoreactivity either due to partial molecular homology between the microbial and host guanosine diphosphate-L-fucose synthase or the genetic constellation of the host. The latter concept seems to be supported by the findings that there are significant associations between the reactivity to GDP-L-fucose synthase peptides along with reactivity against an immunodominant myelin basic protein peptide and the HLA-DRB3*02:02 haplotype, one of the MS-associated genetic constellation found by the International Multiple Sclerosis Genetics Consortium [58].

Interestingly, fucosylated glycans are highly expressed in brain tissue and human CNS myelin compared with other tissues and support recognition by microglia and dendritic cells [182,183]. Thus, one can speculate that the absence of this tolerogenic signal upon reduced fucosylation due to the potentially intestinal microbiome-mediated occurrence of GDP-L-fucose synthase-reactive CD4⁺ T cells contributes to the neuroinflammatory state of the CNS of MS patients.

6. Conclusions

The prevalence of multiple sclerosis has significantly increased over the past three decades, putting a huge economic burden on healthcare systems worldwide [184,185]. The increase in prevalence is attributed to increased incidence, improvement in patient care leading to an increased survival of patients and better diagnostic tools [186,187].

Despite the great efforts made in the research of the disease, details of the pathogenesis remain elusive. Although there is a wide consensus on the autoimmune nature of MS, little is known of the stimuli that trigger the disease or the conditions that make individuals prone to MS. Indeed, although the correlation between certain genotypes or pathogens and the risk of MS has been postulated, the nature of the role of these putative risk factors is still not clear. Advances in the identification of key players in the symptomatic phases of MS, however, could take us closer to answering this question.

Th17 cells have been identified as one of the key players in the pathogenesis of MS. Their hallmark cytokine product, IL-17, however, was found to be protective in both *K. pneumoniae* infections and in asthma models, where it reduced the eosinophilia and bronchial hyper-reactivity alike, suggesting a bone fide role of Th17 cells in these immune situations [188,189]. Moreover, IL-17 was shown to control the integrity of tight junctions in the intestinal epithelium by influencing the subcellular localization of occludin, a critical component of *zonula occludens* [190]. These interactions immediately provoke the question of whether similar microbial-Th17 links give rise to the myelin-reactive populations of Th17 cells via, for instance, molecular mimicry and, if so, whether analogous interplays exist between members of the gastrointestinal microbiome and Th17 lymphocytes.

Indeed, data on the gut microbiome diversity in monozygotic twins discordant to MS, for instance, suggest that the proposed genetic background is not sufficient for the development of the disease, modulating the picture on the importance of genetic risk factors in MS further [153]. Moreover, the microbiomes of MS-twins can provoke autoimmunity accompanied by paralyzed production of the anti-inflammatory IL-10 following transplantation in susceptible animal models, fueling the idea of the pathogenic role of the gut microbiome mediated via immunological elements in MS.

High individual diversity of gut microbiomes may prove the identification of pathogenic species to be particularly difficult, suggesting that research on the role of the human microbiome in diseases like MS should, perhaps, be focused on the characteristics of the protective microbiome composition. Observations that increasing the abundance of *Adlercreutzia* in the gut microbiome of RR-MS patients leads to a decrease in the inflammatory state and consequently delays the progression of MS apparently support this idea, and raise the question of whether therapeutic alteration of the gut microbiome could be useful in the treatment of MS patients.

Indeed, fecal microbiota transplantation (FMT) has been successfully used in diverse conditions including severe dysbiosis due to refractory and recurrent *C. difficile* infections [191]. Although concerns of FMT include the invasiveness of the intervention, the modest but significant risk of infection transmission, thorough donor screening or transplantation of bacteria-free filtrates have already been demonstrated as measures to minimize the risk factors of FMT [192,193]. Following this logic, the use of FMT in MS has been recently tested in a small scale pilot study demonstrating its safety and tolerability, suggesting a new avenue in the treatment of multiple sclerosis [194].

Due to its apparently overwhelming complexity, one can speculate that a great proportion of the mechanism underlying the pathology of MS is highly individual. If that complexity exists, it calls for further investigations to identify mechanisms common in MS patients that could serve as potential targets for developing successful treatment regimes. It is believed that Hippocrates, the forerunner of modern evidence-based medicine, said that “People think that epilepsy is divine simply because they don’t have any idea what causes epilepsy. But I believe that, someday, we will understand what causes epilepsy, and at that moment, we will cease to believe that it’s divine. And so, it is with everything in the universe”. Indeed, life sciences have been on a great journey since the era of Hippocrates

and solved several mysteries woven together with the human body, achievements Hippocrates could not even have dreamed of. Still, we, perhaps, did not pay enough attention to his observation when he said, “All disease starts in the gut”. In the light of advances in the field of intestinal physiology in recent years, it seems that another Hippocrates’ quote, “Let food be thy medicine and medicine be thy food”, perhaps, has never been timelier.

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