Dietary Fatty Acids Directly Impact Central Nervous System Autoimmunity via the Small Intestine

Highlights

- Dietary fatty acids have profound influence on T cell differentiation in the gut
- Middle- and long-chain fatty acids (LCFAs) support Th1 and Th17 cell differentiation
- Short-chain fatty acids (SCFAs) lead to increased Treg cell differentiation
- LCFAs worsen disease in an animal model of MS; SCFAs exert the opposite effect

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In Brief

Haghikia and colleagues show that dietary fatty acids (FAs) influence T cell differentiation in the gut, with short FAs leading to increased Treg cell differentiation and long FAs supporting Th1 and/or Th17 cell differentiation. These FAs differentially affect EAE severity, demonstrating a direct dietary impact on central nervous system autoimmunity.
Dietary Fatty Acids Directly Impact Central Nervous System Autoimmunity via the Small Intestine

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SUMMARY

Growing empirical evidence suggests that nutrition and bacterial metabolites might impact the systemic immune response in the context of disease and autoimmunity. We report that long-chain fatty acids (LCFAs) enhanced differentiation and proliferation of T helper 1 (Th1) and/or Th17 cells and impaired their intestinal sequestration via p38-MAPK pathway. Alternatively, dietary short-chain FAs (SCFAs) expanded gut T regulatory (Treg) cells by suppression of the JNK1 and p38 pathway. We used experimental autoimmune encephalomyelitis (EAE) as a model of T cell-mediated autoimmunity to show that LCFAs consistently decreased SCFAs in the gut and exacerbated disease by expanding pathogenic Th1 and/or Th17 cell populations in the small intestine. Treatment with SCFAs ameliorated EAE and reduced axonal damage via long-lasting imprinting on lamina-propria-derived Treg cells. These data demonstrate a direct dietary impact on intestinal-specific, and subsequently central nervous system-specific, Th cell responses in autoimmunity, and thus might have therapeutic implications for autoimmune diseases such as multiple sclerosis.

INTRODUCTION

Renewed focus on the gut, the largest zone of interaction between the environment and the human organism, has opened new avenues for various fields of life sciences engaged in health and disease. A rapid and simultaneous paradigm shift in microbiologic diagnostics from classical culturing to next-generation sequencing has enabled a more precise estimation of the human gut microbiome composition under healthy conditions (Lozupone et al., 2012). Consequently, there is fast-growing quest for possible disease associations involving interactions between diet, the gut, and microbiome components, especially in autoimmunity (e.g., diabetes and inflammatory bowel disease [IBD]) (Brown et al., 2013; Clemente et al., 2012).

In spite of many remaining questions regarding which components of the microbiome are critically responsible for the fine-tuning of adaptive immune responses in the gut, growing empirical evidence suggests that nutrition and bacterial metabolites might impact the systemic immune response in the context of disease and autoimmunity (Cotillard et al., 2013; Macia et al., 2012). Here, fatty acids (FAs), as an integral component of daily diet, have become a primary focus of investigation. Although the downstream mechanisms and the cellular mediators for the effect of SCFAs remain inconclusive, most data point to the involvement of regulatory immune mechanisms. For example, in models of IBD, gavage of various SCFAs leads to local...
expansion of intestinal T regulatory (Treg) cells (Smith et al., 2013; Furusawa et al., 2013), and microbiome analyses of a large cohort of affected individuals with type 2 diabetes revealed a lack of butyrate-producing bacteria, further underscoring the impact of FAs in health and disease (Qin et al., 2012).

The gut microbiome, along with various dietary habits such as high salt intake, has been recently established as an environmental contributor to the pathogenesis of multiple sclerosis (MS) (Berer et al., 2011; Kleinewietfeld et al., 2013), a T-cell-mediated autoimmune disease of the central nervous system (CNS) with neurodegenerative features (Haghikia et al., 2013). Previous interdisciplinary research has led to the contemporary view that the autoimmune basis of MS stems from an imbalance between pathogenic pro-inflammatory Th1 and/or Th17 cells and anti-inflammatory or regulatory mechanisms of immune cells including Treg cells (Kleinewietfeld and Hafler, 2014). Thus, the rationale of currently available therapeutic interventions is to suppress pathogenic Th1 and Th17 cells and/or to augment Treg cell differentiation (Haghikia et al., 2013).

Here, we show that dietary-induced changes in the gut shaped Th cell responses through the opposing effects of dietary SCFAs and the less-well-studied medium-chain (MC) FAs or LCFAs.

RESULTS

LCFAs Promote Polarization of Naive T Cells toward Th1 and Th17 Cells

To investigate effects of alkanolic acids on the differentiation of naive T cells under Th1 and Th17 cell polarizing conditions in vitro, we tested derivatives with different aliphatic chain lengths in murine CD4+ T cells (Figure 1A and 1B). At 250–500 μM, the C12 FA dodecanoic acid (lauric acid [LA]) increased differentiation of Th17 cells (Figure 1C) and Th1 cells (Figure 1D) by ~50% as compared to control conditions; LA decreased the differentiation of CD4+ T cells from healthy human donors (Figures S1A and S1B). Growth curves and apoptosis rates did not differ between controls and LA-treated cultures, indicating that LA acted to directly enhance T cell differentiation and did not merely affect T cell expansion or viability (Figures S1A–S1L). The addition of LA versus control to naive (CD45RA+CD45RO−CD127+CD25−) CD4+ T cells from healthy human donors also increased the differentiation of CD4+CD25−Foxp3+ Treg cells by about one third versus control (Figure 1E). Expression analyses of differentiated T cell gene signatures revealed increased mRNA expression of Ifng, granulocyte macrophage colony-stimulating factor (Csf2), and tumor necrosis factor alpha (Tnf), but not interleukin-17A (Il17a) for Th1 cells (Figure S2A). Congruent analyses also revealed increased mRNA levels of Il17a, Csf2, Rorc, aryl hydrocarbon acid (lauric acid [LA]) increased differentiation of Th17 cells (Figure 1C) and Th1 cells (Figure 1D) by ~50% as compared to control conditions; LA decreased the differentiation of CD4+ T cells from healthy human donors (Figures S1A and S1B). Growth curves and apoptosis rates did not differ between controls and LA-treated cultures, indicating that LA acted to directly enhance T cell differentiation and did not merely affect T cell expansion or viability (Figures S1A–S1L). The addition of LA versus control to naive (CD45RA+CD45RO−CD127+CD25−) CD4+ T cells from healthy human donors also increased the differentiation of CD4+CD25−Foxp3+ Treg cells by about one third versus control (Figure 1E). Expression analyses of differentiated T cell gene signatures revealed increased mRNA expression of Ifng, granulocyte macrophage colony-stimulating factor (Csf2), and tumor necrosis factor alpha (Tnf), but not interleukin-17A (Il17a) for Th1 cells (Figure S2A). Congruent analyses also revealed increased mRNA levels of Il17a, Csf2, Rorc, aryl hydrocarbon
receptor (Ahr), Tbx21, Ili23a, IL-23 receptor (Il23r), transforming growth factor beta (Tgfb1), and Tnf, but not Ifng, in the presence of 250 μM LA versus control (Figure 2B), supporting the argument for an increased generation of Th17 cells after addition of LA in vitro. Treg cell quantitative real-time PCR (qRT-PCR) revealed a ~50% reduction of Foxp3 expression after addition of LA versus control in vitro (Figure 2C).

Gene and Protein Expression Analyses Implicate Th17 Cell and MAPK Pathways as Modulators of LA Cellular Mechanism

To further understand the differential effects of different FA lengths on T cell differentiation, we analyzed gene expression of the candidate fatty acid receptors liver X receptor alpha (LXRa) and various G protein coupled receptors (GPR40, GPR41, GPR43, GPR84, GPR119, and GPR120) in different CD4+ T cell subsets. The presence of GPR40, 41, 43, and 84 on the mRNA level was mostly confined to naive T cells. In contrast to previous reports (Smith et al., 2013), none of the receptors were detectable in Th17 cells or Treg cells, and only Gpr43 was present in Th1 cells. Lxra, Gpr119, and Gpr120 were not detected in any T cell subset (Figure 2D). Likewise, addition of the Toll-like receptor 1/2 agonist Pam3CSK4 to Th1, Th17, or Treg cell differentiation assays did not show any comparable effect to LA (not shown).

No additional increase of Th17 cell frequency was found subsequent to the addition of LA to dendritic cells (DCs) cocultured with LA-treated T cells, arguing for a direct effect of LA in fostering Th17 cell polarization (Figure 2A). In order to identify downstream signaling factors potentially involved in the LA effect on Th17 cell differentiation, we evaluated the transcriptome profile of in vitro, LA-treated versus -untreated Th1 and Th17 cells. Analyses of Th17 cells revealed Maf (c-Maf) as the highest differentially expressed gene (DEG), increasing 16.5-fold in LA- versus control-treated T cells (Figure 2B). Further DEGs pointing to induction of Th17 cell differentiation in multiple pathway analyses included the salt-sensing kinase Sgk1 and multiple MAPK family members, such as Mapk14 (encodes p38). The expression profile of Th1 cells revealed differential expression of critical Th1 cell genes, whereas changes in the MAPK family were not as prominent (not shown).

Because p38 MAPK-mediated signaling is a well-characterized integrator of environmental stressors, we tested the effects of FAs on p38 MAPK regulation. Expression analysis revealed a statistically significant ~13-fold increase in Mapk14 in Th17 cells, and to a lesser degree in Th1 cells, upon LA application (Figure 2C). At the post-transcriptional level, LA treatment of T cells under Th17-cell-polarizing conditions led to a significant increase of p38 phosphorylation (p38, Figure 2D). Pharmacological inhibition of p38 MAPK via addition of the specific blocker SB202190 almost completely inhibited the effect of LA on Th17 or Th1 cell polarization (Figures 2E and 2F). Furthermore, genetic ablation of the p38α subunit via CD4-cre-mediated deletion in T cells completely abolished the enhanced Th17 and Th1 cell differentiation effect of LA (Figures 2G and 2H). Neither pharmacological inhibition nor genetic ablation of p38 led to increased cell death (not shown).

SCFAs Promote Polarization of Naive T Cells toward Treg Cells

Corresponding to the effect of FAs on Th1 and Th17 cell differentiation, we tested derivatives with different aliphatic chain lengths on naive T cells under Treg-cell-polarizing conditions (Figures 3A and 3B). Propionate (PA, C3:0) revealed the most significant effect on murine Treg cell differentiation in vitro at a concentration of 150 μM (Figure 3C). We further corroborated the Treg-cell-stimulating effect of PA in vitro in a human T cell differentiation assay. Therein, application of PA to healthy donor naive CD4+ T cells increased both the frequency of CD4+CD25+ Foxp3+ cells and, to a lesser extent, the proliferation of differentiated Treg cells (Figure 3D). Adding PA to human naive T cells under Th17-cell-polarizing conditions led to a significant reduction in the frequency of CD4+IL-17A+ T cells (Figure 3E).

Furthermore, ex vivo transcriptome analyses of Treg cells derived from PA-pre-treated mice and Treg cells from previously untreated mice revealed DEGs in the two different cell populations (Figure 3F). Through the use of multiple pathway analyses steps, rationally selected DEGs revealed, and qRT-PCR confirmed (Figure 3G), Lpin-2 (encodes lipin2) and Mapk8 interacting protein-2 (Mapk8ip2, encodes JIP2) as relevant, upregulated genes in Treg cells from PA-pre-treated mice. In contrast, LA treatment led to a downregulation of Lpin-2 expression in both Th1 and Treg cells; however, changes in Mapk8ip2 were not significant in both cell types. Furthermore, LA treatment had no effect on this pathway in Th17 cells. As previously shown, the increase of Lpin-2 led downstream to downregulation of JNK1 activation (encoded by Mapk8), which in turn is a key activator of pro-inflammatory transcription factors such as NF-κB. Relative mRNA expression of Mapk8 parallels that of Lpin-2 in all investigated cell types after LA application (Figures S3A–S3C). At the post-transcriptional level, PA treatment of T cells led to an opposing effect on p38 phosphorylation (p38, Figure 3H) compared to that which was exerted by LA on Th17 cells (Figure 2D).

LA-Rich Diet Impacts Th1-Cell- and Th17-Cell-Mediated CNS Autoimmunity in Vivo

The distinct in vitro effects of FAs on naive CD4+ cells prompted us to examine their effects in vivo using murine MOG35-55 EAE as a model of Th1-cell- and/or Th17-cell-mediated autoimmunity. Male C57BL/6 mice were fed standardized and otherwise completely matched diets rich in either LA (C12) or palmitic acid (PALM, C16) and were compared to mice on a control diet after EAE induction. Mice on the LA-rich diet did not display different body weights compared to controls (mean ± SEM on day 10 p.i. = 23.2 ± 0.5 g versus 23.8 ± 2.0 g, p = 0.96). Mice fed the LA-/PALM-rich diets displayed a more severe course of the disease (Figures 4A and S4A–S4C), although disease incidence (12/13 ctrl. versus 10/10 LA) and mortality were unaffected. Upon ex vivo phenotyping by flow cytometry of the spinal cord infiltrates, the LA diet increased Th17 cell frequencies in the CNS on day 14 post immunization (p.i.) whereas Th1 cells (Figure 4B) and CD11b+ antigen-presenting cells (Figure 4C) remained unchanged.

Phenotyping of splenic T cells during EAE revealed an increase of Th1 and Th17 cell frequencies under an LA-rich diet on day 10 p.i. (Figure 4D). In parallel, an LA-rich diet expanded immunity 43, 817–829, October 20, 2015 ©2015 Elsevier Inc. 819
Figure 2. MAPK Pathways Are Modulators of LA Cellular Mechanism

(A) Addition of LA to co-culture assay of T cells and dendritic cells generated in the presence (LA-treated BMDCs) or absence (untreated BMDCs) of 250 μM LA (n = 3, one out of three independent experiments shown).

(B) Microarray analysis of in-vitro-generated Th17 cells in the presence (Th17+LA) or absence (Th17 ctrl) of 250 μM LA. Left: Heat map analysis displaying differentially expressed genes (DEG). Right: a selection of 32 up- and downregulated genes.

(C) Gene expression analysis of Mapk14 (p38 MAPK) expression in Th1, Th17, and Treg cell differentiation assays (triplicates, data pooled from two preparations).

(D) Immunoblot analysis of p38 protein and phosphorylated p38 protein (pp38) in Th1 and Th17 cell differentiation assays with and without LA. Left: quantification of p38 and pp38 protein. Right: representative blots. β-actin (β-act) was used as loading control.

(E and F) Chemical inhibition of p38 via the addition of SB202190 to murine CD4+ T cell differentiation culture under Th17 (E) and Th1 (F) cell polarizing conditions (n = 4–7 per group, data pooled from two experiments).

(G and H) Conditional genetic deletion of p38α in T cells via Cre-loxP in a murine T cell differentiation under Th17 (G) and Th1 (H) cell polarization (n = 6 per group, data pooled from two experiments).

*p < 0.05, **p < 0.01; ***p < 0.001. See also Figure S2.
effector T cells on day 14 p.i. with significantly decreased
CD4+CD25+ cell frequency (and increased CD69+ cell frequency, not shown) and increased CD4+CD44+ cell frequency and CD4+CD25+ cell frequency (Figure 4E). Splenocyte recall assays also revealed increased IFN-γ and IL-17A secretion on day 10 p.i. (Figures S4D and S4E). Via the same procedure, LA-rich diet increased T cell proliferation without additional stimulation; however, this effect was more significant in CD4+ and CD8+ T lymphocytes after MOG35-55-specific recall (by 100%) or polyclonal activation by concanavalin A (by >300%, Figure 4F).

LA Exerts Effect on Th17 Cells via the Small Intestine
In order to elucidate the anatomic site of the observed in vivo actions of LA, we examined LA concentrations in the serum, blood cells, the complete duodenum, and duodenal mucosa after either LA-rich or control diet on day 10 p.i. We found that LA concentrations in the blood compartment (Figure 5A), mucosa, and duodenum (Figure 5B) were significantly increased after LA diet, with the highest concentrations in the blood compartment and an enrichment of LA in the duodenal mucosa versus the complete duodenum. More extensive analysis of FAs in these compartments also revealed an increase of MCFAs and LCFAs of various chain lengths, especially C14–C18 under an LA diet (Figures S5A–S5D). Given the enrichment of LA in the small intestine (SI) mucosa, we further scrutinized the gut. In a kinetic analysis of intraepithelial lymphocytes (IELs), lamina propria lymphocytes (LPLs), and Peyer’s patches (PPs) in various gut locations, we found the most pronounced increase in IL-17A and IFN-γ on day 10 p.i. with MOG 35-55 versus naive mice. Although EAE itself did not alter microbiome composition, the LA diet reduced Prevotellaceae and S24-7 families of the Bacteroidetes phylum as compared to controls (Figure 5I and Table S1).

In Vivo, PA-Mediated Treg Cell Regulatory Response Ameliorates CNS Autoimmunity
To evaluate whether the opposing effect of the C3 FA PA seen in vitro exerts an ameliorating effect on the course of EAE, we applied 150 mM PA (or water solvent) by daily oral gavage either (1) at the day of immunization (DI) or (2) at onset of disease (OD). In contrast to the LA-rich diet, application of PA revealed beneficial effects in EAE, but only in the preventive setting (Figure 6A). Ex vivo recall assays revealed a significant increase of the anti-inflammatory cytokine IL-10 in cells derived from PA-treated EAE, whereas, in contrast to LA-diet mice, no significant difference in IL-17A production was observed (Figure 6B). Phenotyping of SI LPLs on day 8 p.i. revealed increased CD4+CD25+Foxp3+ Treg cell frequency in PA-treated EAE mice (Figure 6C). Analyses of the signature cytokines in different gut sub-compartments also showed increased mRNA levels of Tgfb1, Il10, and Foxp3 in the distal parts of the SI in these mice (Figure 6D). To test whether orally delivered PA imprints a protective phenotype on Treg cells in vivo, we transferred Treg cells derived from PA-pre-treated (7 days) or untreated (5 × 10^5 CD4+CD25+ Treg cells; i.p.) mice into recipient mice concurrently with EAE induction; flow cytometry analysis of isolated cells confirmed the congruity of Foxp3+ cell percentages in the transferred T cell preparations (69%–72% under both conditions). Treg cells derived from PA-pre-treated mice led to a marked improvement in the clinical course of the recipient EAE as compared to controls (Figure 6E). The improved clinical outcome in PA-DI-treated mice was reflected in the histopathological analyses of the spinal cord, with less inflammatory cell infiltration, less demyelination, and a higher degree of axonal preservation (Figure 6F).

DISCUSSION
Our data add FAs to the list of environmental triggers of T cell differentiation that might act as risk factors for Th1-cell- and/or...
Figure 3. SCFAs Promote Polarization of Naive T Cells toward Treg Cells and Suppress the JNK1 and p38 Pathway

(A) Addition of PA derivatives to murine CD4+ T cells under Treg cell polarizing conditions. C3:0 PA, C4:0 butyric acid, C6:0 caproic acid, C8:0 caprylic acid, C10:0 capric acid, C12:0 lauric acid (all FAs at 100 μM; n = 7).

(B) Addition of different PA concentrations to murine Treg cell differentiation (n = 8).

(C) Addition of PA to murine CD4+ T cell differentiation culture under Treg cell polarizing conditions (n = 8).

(D) Addition of PA to human CD4+ T cell proliferation and differentiation culture under Treg cell polarizing conditions (n = 5, one out of two experiments shown).

(E) Addition of PA to human CD4+ T cell proliferation and differentiation culture under Th17 cell polarizing conditions (n = 5, one out of two experiments shown).

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Th17-cell-mediated autoimmune diseases like MS (Kleinewietfeld and Hafler, 2014). Recent epidemiological studies reveal that lifestyle factors such as smoking, obesity, and salt intake might constitute a risk for MS (Hedström et al., 2011, 2014; Kleinewietfeld et al., 2013). Because such a diet is also typically rich in FAs, our present data offer important immunological and functional groundwork for epidemiological observations and identify saturated FAs as a new dietary, non-infectious trigger involved in (F) heat map analysis displaying differentially expressed genes (DEG) in the transcriptome analysis of Treg cell derived from PA pre-treated or previously untreated (control) mice (left). Right: a selection of 24 up- and downregulated genes.

Figure 4. LA Exacerbates CNS Autoimmunity and Increases Th1 and Th17 Cells In Vivo
(A) Clinical course of MOG35-55 EAE. Mice were fed a LA-rich diet (n = 10) or control diet (n = 13) for 4 weeks prior to immunization. Data are shown on a 5-point score scale pooled from two experiments. (B and C) Ex vivo flow cytometry analysis of (B) Th1 and Th17 cell frequencies and CD11b+ cells (C) in the spinal cord under LA-rich versus control diet on day 14 of MOG35-55 EAE (n = 7 per group). (D) Ex vivo flow cytometry analysis of Th1 and Th17 cell frequencies in the spleen on day 10 of MOG35-55 EAE (n = 7 versus 10 mice per group). (E) Ex vivo flow cytometry analysis of CD25+, CD44+, or CD62L+ effector T cells from spleen on day 14 of MOG35-55 EAE under LA-rich versus control diet (n = 7 per group). (F) Proliferation of CD4+ and CD8+ T cells after ex vivo recall with MOG35-55 (splenocytes harvested on day 14 p.i. of MOG35-55 EAE, n = 4 per group). Ø = none ns = not significant, *p < 0.05, **p < 0.01; ***p < 0.001. See also Figure S4.

intestinal T cell differentiation due to the interaction of gut-hosted microbiota and nutritional metabolites.

Here we show that, in autoimmune disease, FAs exert direct effects both on murine and human Th cells under polarizing conditions in vitro and in the context of antigen-specific T cell response in vivo. Treatment with SCFAs, PA most potently, enhanced differentiation and proliferation of CD4+CD25+Foxp3+ Treg cells and ameliorated EAE disease course. In contrast, MCFAs or LCFAs such as LA or PALM enhanced Th1 and Th17 cell differentiation and contributed to a more severe course of EAE. The simultaneous increase of tracked Th17 cells in the spleen and CNS together support a model of differential segregation of antigen-specific T cells under an LA-rich diet. This hypothesis is further supported by the increased levels of LA found in the blood and mucosa compartments in mice having received an LA-rich diet. The increase of LA in both compartments might explain not only the systemic effect of LA on Th17 cell differentiation in vivo, but also the increased frequency of Th17 cells in the lamina propria during the initial antigen-specific immune response in EAE.

(F) Heat map analysis displaying differentially expressed genes (DEG) in the transcriptome analysis of Treg cell derived from PA pre-treated or previously untreated (control) mice (left). Right: a selection of 24 up- and downregulated genes.

(G) Validation of DEG detected by the transcriptome analysis via qRT-PCR showing upregulation of Lpin-2 (lipin-2) and marked mRNA upregulation of mapk8ip2 (JNK1 interacting protein2, JIP2) in Treg cells derived from mice pre-treated with PA as compared to Treg cells from untreated mice (from five independent preparations).

(H) Immunoblot analysis of p38 and phosphorylated p38 protein (pp38) in Treg cell differentiation assays with and without PA. β-actin (β-act) was used as loading control.

*p < 0.05, **p < 0.01, ***p < 0.001. See also Figure S3.
The proposed FA-receptor mediating the aforementioned effects remains to be identified. The presented divergent effects of saturated FAs on Th1 and Th17 versus Treg cells depended on the length of the carbon chain backbone, thus arguing for a pattern- or chain length-sensing receptor on T cells. Here, GPR109 might be a critical player, as has been recently shown for butyric acid (Singh et al., 2014), whereas receptors previously noted of importance, GPR41 and GPR43 (Kim et al., 2013), are not as relevant in our system. Given the plasticity of Th17 and Treg cells and their complex, divergent, intercellular signaling pathways during differentiation (for review see Kleineveldt and Hafler, 2014), it is likely that several mechanisms act in concert.

On both the gene expression and post-transcriptional levels (protein), members of the MAP kinase family, including p38 and JNK1, along with lipin-2 and their regulation of downstream pathways, are herein identified as crucial mediators of the effects of saturated FAs on T cells. Although both p38 and JNK1 have been shown to non-redundantly contribute to T cell death, differentiation, and proliferation (Rincón and Pedraza-Alva, 2003), p38-MAPK in particular is a well-known integrator of environmental stress and is involved in both T cell differentiation (Kleineveldt et al., 2013) and models of MS (Kremensov et al., 2014; Noubade et al., 2011). Because p38-MAPK plays a prominent role in T cell development and function (Alam et al., 2014), it can be argued that developmental changes due to the genetic knock-out of p38-MAPK lead to future alterations of T cell differentiation prior to the application of FAs. However, pharmacological blockage of this pathway after T cell development confirmed its specific involvement in FA effects on T cell differentiation. Indeed, the coupling of both the p38 and JNK MAP kinase pathways to T cell receptor signaling might allow for lineage-specific signals in T cell differentiation (Flavell et al., 1999). Additionally, our identification of lipin-2 signaling in the PA-induced generation of Treg cells adds to previous literature whereby it has been shown to counteract the pro-inflammatory effects of saturated FAs on macrophages (Valdearcos et al., 2012), yet it had not been previously described in the context of T cells.

In addition to the MAP kinase family members, transcription factor c-Maf and kinase SGK1, both of which have been shown to be critically involved in Th17 cell differentiation (Tanaka et al., 2014; Wu et al., 2013), were among the differentially expressed genes identified in LA-treated T cells under Th17-cell-polarizing conditions.

In the absence of antigen, the expansion of Th17 cells in the SI of anti-CD3-treated mice clearly increased under LA feeding. Despite its limitations, especially in the context of CNS autoimmunity, the ability of anti-CD3 to induce acute immune activation offers a valuable setup to shed further light on the distribution of auto-antigen-specific Th cells under an LA-rich diet (Esplugues et al., 2011). Indeed, our observation of an increased influx of Th17 cells into the SI, e.g., the lamina propria, of LA-fed mice during EAE, provides evidence for the involvement of the SI, including nutritional metabolites and microbiota of the gut, in systemic T cell immune responses prior to T cell CNS infiltration.

The shift of the gut microbiome toward decreased Prevotellaceae and Bacteroidetes after an LA-rich diet might partially explain the enhanced positive effect of fecal transplantations over diet manipulation. This concept is backed by the observed decrease in SCFAs in the gut under LA-rich diet along with recent reports implicating the pivotal role of Bacteroidetes in the fermentation of fiber-rich nutrition into SCFAs. Furthermore, a dysbiotic microbiome with lower proportions of Bacteroidetes has been associated with immune dysregulation and incidence of autoimmune disorders such as Crohn’s disease and asthma (Macia et al., 2012; Trompette et al., 2014).

Of particular interest is the ability of PA to beneficially influence the generation of Treg cells (Arpaia et al., 2013). Our study posits PA as a potent compound with the capacity to restrain CNS autoimmunity via restoration of the altered Treg cell:effector T cell balance, which is disturbed in MS patients (Viglietta et al., 2004). So far, influencing MS via direct Treg cell manipulations, e.g., via superagonistic anti-CD28 antibodies, has been considered worthwhile (Beyersdorf et al., 2005) but not practically feasible outside of controlled experimental conditions (Hüning, 2012). In fact, PA is a common environmental compound that was traditionally part of many preservatives and is already ingested by many people at lower concentrations (Cummings et al., 1987). Given this wide range in everyday practice and ingestion, it is likely that several mechanisms act in concert.
Figure 6. PA Ameliorates CNS Autoimmunity via Induction of Treg Cells in the Small Intestine

(A) Clinical course of MOG35-55 EAE. Mice received daily PA starting either at the day of immunization (DI, n = 15), at onset of disease (OD, n = 25), or the solvent water (controls, n = 29) via oral gavage in addition to the normal diet. Data are shown on a 5-point score scale pooled from three experiments.

(B) Cytokine measurement in cultures after ex vivo recall with MOG35-55 (splenocytes harvested on day 8 p.i. of MOG35-55 EAE, n = 8 versus 9 per group).

(C) Flow cytometry analysis of cells isolated from the LPLs of EAE either treated with PA (n = 5) or with water (untreated, n = 5) showing higher amount of T reg cells in PA-treated mice.

(D) Gene expression analyses of tgfβ1, il10, and foxp3 in the SI LPLs of either PA (PA DI) or water (control) (triplicates, data pooled from seven to ten preparations).

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growing evidence for the lasting dietary effect on microbiome composition (Cottillard et al., 2013), a rapid translation of PA therapy from pre-clinical studies to clinical trials in MS patients seems reasonable.

In sum, this study identifies dietary saturated FAs as crucial modulators in the gut, shifting Th1 and Th17 versus Treg cell balance in autoimmune neuroinflammation. We propose a dual mechanism of action for LA, comprising (1) increased Th17 and Th1 cell polarization and proliferation as a systemic effect and (2) enhanced sequestration of Th17 cells in the gut in the context of immune activation. Further studies in humans to explore the supplementary therapeutic potential of enriched diets are highly warranted.

**EXPERIMENTAL PROCEDURES**

**EAE and Diet**
For experiments under LA-rich diet, mice received a chow containing 30.9% crude fat rich in the MCFA C12:0 (lauric acid, 13.47%), and mice on a control diet were fed a chow with 4.2% crude fat. Mice were adapted to high-fat chow for 4 weeks before EAE induction. For EAE under LA, either 200 μl PA (150 mM) or solvent (water) were applied daily via oral gavage in addition to normal diet with 3.3% crude fat, at either day of immunization (DI) or onset of disease (OD). For transfer experiments, mice received either PA or water via oral gavage in addition to normal diet for 7 days before gut Treg cells were isolated and injected into EAE at DI.

For EAE, 8- to 11-week-old mice were anesthetized and subcutaneously injected with 200 μg MOG35-55, and 200 μg CFA. Pertussis toxin (200 ng/mouse) was applied i.p. on days 0 and 2 p.i. Daily clinical evaluation was performed via a 5-point scale. All experiments were performed in accordance with the German laws for animal protection and were approved by the local ethic committees (Erlangen AZ 54-2532.1-56/12; Bochum AZ 84-02.04.2014.A104). For details and providers of materials used, see **Supplemental Information**.

**In Vitro MOG Restimulation Assay**
Splenocytes from EAE mice were obtained on day 10 p.i. and seeded at a density of 3 x 10^5 cells/ml in Re-medium. MOG35-55 (1, 20, 100 μg/ml) and ConA (1.25 μg/ml) were added for stimulation, and cells were cultured 48 hr. Supernatants were harvested and analyzed for cytokines (ELISA). To monitor proliferation, cells were labeled with fixable proliferation dye (eBioscience) according to the manufacturer’s protocol.

**Cell Tracking Experiment**
Splenocytes from EAE mice were obtained on day 10 p.i., labeled with fixable e450 proliferation dye (eBioscience), and intravenously transferred into mice on a LA or control diet on day 10 p.i. (25 x 10^6 cells/mouse). Splenocytes, CNS, and SI of recipient mice were scrutinized on day 4 post cell transfer and analyzed by flow cytometry for CD4^+e450^ cells.

**Free Fatty Acid Quantification**
Total fatty acid concentrations in plasma, whole blood cells, and duodenum of mice on a LA or control diet were analyzed on day 10 p.i. via gas chromatography (GC) as described (Osternann et al., 2014). In brief, lipids were extracted with MTBE/MeOH and derivatized with methanolic hydrochloric acid, and the resulting FAME were quantified by GC with flame ionization detection. For details of SCFAs analyses in feces, see **Supplemental Information**.

**In Vivo MOG Restimulation Assay**
Splenocytes from EAE mice were obtained on day 10 p.i., injected with anti-CD3 (20 μg, 145-2C11, BD PharMingen) or with PBS alone or CFA in PBS and sacrificed after 3 days.

**Human T Cell Differentiation**
PBMCs from whole blood of healthy volunteers were separated by Ficoll-Paque PLUS gradient centrifugation and naïve (CD45RA^+CD62L^+CD44^loCD25^-) were stimulated by 2 ng/ml anti-CD3 and 1 ng/ml IL-2 (1 ng/ml); IL-1 (10 ng/ml); IL-6 (25 ng/ml); IL-1β, IL-23 (25 ng/ml), and rh-TGF-β1 (10 ng/ml) for Th17 cells; rh-TGF-β1 (5 ng/ml) for Treg cells. For investigating the influence of FAs on T cell differentiation, PALM and LA were added to the cultures. All cells were analyzed via flow cytometry analysis. For details and providers, see **Supplemental Information**.
Fecal Transplant

200 mg feces from mice on LA or control diet were collected in a sterile manner on the day of fecal transfer and re-suspended in 1 ml sterile filtered autoclaved tap water and adjusted to a final volume of 200 mg feces/ml, vortexed, and centrifuged. Each recipient mouse received 200 μl of fecal supernatant by oral gavage on 3 consecutive days before αCD3 injection (20 μg/mouse, 145-2C11, BD Pharmingen) after 3 weeks of housing in sterile cages with sterile autoclaved water and autoclaved food.

Transcriptome Analysis

Total RNA was isolated from corresponding cells using the RNaseasy Mini Kit (Qiagen) with prior pepGold Trifast (Peqlab) treatment. cDNA was generated with Ambion WT Expression Kit (Life Technologies). Subsequently, Affymetrix GeneChip Mouse Gene ST Arrays were processed by the manufacturer’s protocol. Data were processed with the Affymetrix Expression Console and Transcriptome Analysis Console v.2.0 (TAC) software, and candidate genes were selected rationally under consideration of both gene functions found in literature (NCBI and GeneCards) and integrative pathway analyses by Ingenuity Pathway Analysis (IPA, Qiagen).

Statistical Analysis

Statistical analysis was performed with GraphPad Prism (GraphPad Software). All in vitro and ex vivo data were analyzed by one-way ANOVA followed by Tukey’s post test, unpaired t test, or Wilcoxon rank sum test after checking for normal distribution (unless indicated otherwise in the legends). EAE data were processed with GraphPad Prism (GraphPad Software). Statistical analysis was performed with GraphPad Prism (GraphPad Software). Statistical analysis was performed with GraphPad Prism (GraphPad Software).

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