The Short Chain Fatty Acid Butyrate Imprints an Antimicrobial Program in Macrophages

**Highlights**

- Butyrate induces differentiation of macrophages with potent antimicrobial function
- Enhanced antimicrobial function is a consequence of glycolysis and mTOR inhibition
- Single-cell RNA-sequencing identifies butyrate-induced antimicrobial peptides
- Butyrate inhibits HDAC3 to drive metabolic changes and microbicidal function

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**In Brief**

Macrophages maintain gut homeostasis by eliminating invasive pathogens and regulating inflammatory responses. Schulthess et al. demonstrate that butyrate, a bacterial fermentation product, imprints potent antimicrobial activity during macrophage differentiation through HDAC3i function.

Schulthess et al., 2019, Immunity 50, 432–445
February 19, 2019 © 2018 The Authors. Published by Elsevier Inc.
https://doi.org/10.1016/j.immuni.2018.12.018
The Short Chain Fatty Acid Butyrate Imprints an Antimicrobial Program in Macrophages

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https://doi.org/10.1016/j.immuni.2018.12.018

SUMMARY

Host microbial cross-talk is essential to maintain intestinal homeostasis. However, maladaptation of this response through microbial dysbiosis or defective host defense toward invasive intestinal bacteria can result in chronic inflammation. We have shown that macrophages differentiated in the presence of the bacterial metabolite butyrate display enhanced antimicrobial activity. Butyrate-induced antimicrobial activity was associated with a shift in macrophage metabolism, a reduction in mTOR kinase activity, increased LC3-associated host defense and antimicrobial peptide production in the absence of an increased inflammatory cytokine response. Butyrate drove this monocye to macrophage differentiation program through histone deacetylase 3 (HDAC3) inhibition. Administration of butyrate induced antimicrobial activity in intestinal macrophages in vivo and increased resistance to enteropathogens. Our data suggest that (1) increased intestinal butyrate might represent a strategy to bolster host defense without tissue damaging inflammation and (2) that pharmacological HDAC3 inhibition might drive selective macrophage functions toward antimicrobial host defense.

INTRODUCTION

The gastrointestinal tract is colonized by a high density of commensal bacteria and is a major site of pathogen entry (Rooks and Garrett, 2016) requiring robust barrier function. Short chain fatty acids (SCFAs) are derived from bacterial fermentation of dietary fibers in the colonic lumen. The SCFAs butyrate, propionate, and acetate promote intestinal epithelial barrier function and regulate the host mucosal immune system (Vinolo et al., 2011b). For example, butyrate serves as a primary energy source for intestinal epithelial cells, the first line of cellular defense against invading pathogens. Butyrate also regulates stem cell turnover in intestinal epithelial crypts (Kaiko et al., 2016). SCFAs, and in particular butyrate also promote regulatory T cells (Treg) in the colon by inhibiting histone deacetylase (HDAC) activity at the Foxp3 locus (Arpaia et al., 2013; Furusawa et al., 2013; Smith et al., 2013). Furthermore, exposure of peripheral blood mononuclear cells such as neutrophils, macrophages, and dendritic cells to SCFAs or other HDAC inhibitors, such as trichostatin (TSA), inhibits inflammatory cytokine production (Chang et al., 2014; Usami et al., 2008; Vinolo et al., 2011a).

Mouse models of intestinal inflammation suggest that butyrate plays an immune regulatory role in vivo (Furusawa et al., 2013). This is potentially relevant for human immunopathology since reduced numbers of butyrate-producing bacteria were found in the gut mucosa and in fecal samples from patients with inflammatory bowel disease (IBD) or colon cancer (Frank et al., 2007; Wang et al., 2012).

Intestinal phagocytes, and tissue-resident macrophages in particular, act as an innate barrier in the intestine by clearing invading bacteria. Malfunctioning of this pathway is involved in the pathogenesis of IBD since defective microbiocidal responses were identified in polygenic and monogenic forms of IBD (Peloquin et al., 2016; Uhlig and Powrie, 2018). In contrast to macrophages found in other organs, intestinal macrophages are largely replenished from blood monocytes (Bain et al., 2014). Thus, circulating monocytes enter the gut and undergo final differentiation in the lamina propria to become mature, highly phagocytic macrophages capable of bactericidal activity via mechanisms such as NADPH-oxidase-derived reactive oxygen species (ROS) and antimicrobial peptides and proteins (Bain et al., 2014; Smythies et al., 2005; Varol et al., 2009). The bacterial pathways that shape macrophage host defense in the intestine are poorly understood. Here we have investigated the ability of SCFAs to influence macrophage function. We show that SCFAs induce metabolic and transcriptional changes in macrophages, which enhances their bactericidal functions.
RESULTS

Butyrate Exposure during Macrophage Differentiation Enhances Antimicrobial Activity

To assess the impact of SCFAs on human macrophages, we differentiated peripheral blood-derived CD14+ monocytes with macrophage colony-stimulating factor (M-CSF) in the absence (control macrophages) or presence of butyrate (butyrate macrophages), propionate (propionate macrophages), or acetate (acetate macrophages). The presence of SCFAs during macrophage differentiation did not affect key macrophage characteristics such as morphology and surface expression of CD11c and HLA-DR (Figures S1A and S1B). However, SCFAs did affect the antimicrobial function of macrophages assessed in a gentamicin protection assay using a range of bacteria including gram negative (Salmonella enterica serovar Typhimurium, later on referred to as Salmonella); Crohn’s disease-associated adherent-invasive Escherichia coli (AIEC), and Citrobacter rodentium (C. rodentium) and gram-positive Staphylococcus aureus (S. aureus) microbes (Figures 1A–1D and Figure S1B). As the anti-microbial effect was strongest with butyrate as opposed to proprionate and not observed with acetate, butyrate was studied further (Figure 1E and Figures S1C–S1E).

We next investigated the kinetics of antibacterial activity, cytokine secretion, phagocytic capacity, and apoptosis in control macrophages and butyrate macrophages. Butyrate macrophages displayed significantly increased elimination of intracellular Salmonella as early as 30 min after infection, and this was maintained over the 3 h duration of the gentamicin protection assay (Figure 1F). Despite altered bacterial clearance, control macrophages and butyrate macrophages displayed similar expression of IL1B and TNF mRNA and protein following 3 h of Salmonella infection (Figures S1F and S1G), whereas IL10 expression was reduced in butyrate macrophages (Figure S1H). In contrast to the long-term exposure of butyrate during the differentiation process, incubation of control macrophages with butyrate for 3 h prior to the gentamicin protection assay did not induce a significant reduction in CFU compared to untreated.
Figure 2. Increase of Antimicrobial LC3-Associated Immune Defense in Butyrate Macrophages

(A) Extracellular acidification rate (ECAR) measured at steady state in control and butyrate macrophages. Data represent the mean of nine biological replicates from three independent experiments.

(legend continued on next page)
control macrophages (Figure 1G). After 24 h incubation in butyrate-free media, butyrate macrophages retained enhanced antimicrobial activity, indicating that this response does not require butyrate at the time of infection (Figure 1H). The reduced bacterial load was not due to reduced phagocytosis as butyrate macrophages showed normal ingestion of FITC beads coated with human IgG and uptake of non-opsonised GFP-Salmonella over 90 min. In addition, they showed similar mRNA expression of the phagocytic receptors MARCO and CLEC7A compared to control (Figures S11–S1L). Reduced numbers of bacteria in butyrate macrophages was not a consequence of increased cell death as the percentage of apoptotic cells was also similar between butyrate macrophages and control macrophages (Figures S1M and S1N). Taken together, these results show that the presence of butyrate during the differentiation of macrophages induces long-lasting antimicrobial activity without affecting phagocytosis, inflammatory cytokine production, or apoptosis.

**Butyrate Alters Metabolism and Induces mTOR Dependent LC3-Associated Antimicrobial Clearance in Macrophages**

As butyrate is a key energy source for epithelial cells (Vinolo et al., 2011b) and relevant amounts of butyrate are likely present in the lamina propria as suggested by portal vein concentrations (van der Beek et al., 2015), we tested whether butyrate exposure alters macrophage metabolism. Butyrate macrophages showed a decreased extracellular acidification rate (ECAR) in comparison to control macrophages (Figure 2A). Glycolysis, glycolytic capacity and glycolytic reserve were also significantly reduced in butyrate macrophages (Figures 2B–2D). To explore the metabolic pathways modulated by butyrate, we performed a metabolomic analysis by liquid chromatography–mass spectrometry. This confirmed substantial changes in the glycolysis pathway including a reduced glucose concentration (Figure 2E and Figure S2A). This is unlikely due to reduced glucose uptake as 2NBDG uptake in control and butyrate macrophages was similar (Figure S2B). Since we found similar mitochondrial oxidative phosphorylation (respiration, proton leak, oxygen consumption rate) between control and butyrate macrophages, the switch in mitochondrial metabolism was not compensated for by increased mitochondrial energy metabolism (Figures S2C–S2H).

We found 34 significantly differentially expressed metabolites between control and butyrate macrophages (Figure 2E). Butyrate was the most differentially expressed metabolite in butyrate macrophages confirming the validity of this approach. Butyrate macrophages also contained higher amounts of adenosine monophosphate (AMP) a known inducer of AMP kinase (AMPK) that inhibits mTOR (Inoki et al., 2003) the master regulator of autophagy (Kim and Guan, 2015). In agreement with this hypothesis, AMPK Thr172 phosphorylation was significantly increased in butyrate macrophages compared to controls (Figure 2F). To test whether butyrate treatment blocked mTOR activity, we analyzed ribosomal protein S6 kinase phosphorylation by flow cytometry and immunoblot as a surrogate marker of mTOR activation. Indeed, butyrate macrophages showed a marked reduction of pS6 phosphorylation compared to control macrophages (Figure 2G). In support of the hypothesis that butyrate-induced antibacterial clearance depends on mTOR inhibition, we found that treatment with MHY1485, an mTOR activator (Figure 2H) blunted butyrate macrophage antibacterial activity (Choi et al., 2012).

Since mTOR is a key regulator of autophagy and autophagy-related processes, we tested whether butyrate-induced antimicrobial activity was associated with bacterial-associated autophagy protein microtubule-associated protein 1 light chain 3 alpha (LC3). We tested the LC3-I to LC3-II conversion by immunoblot, LC3 flow cytometry and confocal microscopy. Immunoblot indicated that the basal and Salmonella infection-induced LC3-II turnover was significantly increased in butyrate macrophages (Figure 2I and 2J). In contrast to the LC3-II increase, P62 protein (SQSTM1) expression was unchanged between control and butyrate macrophages (Figure 2I and 2K), indicating an overall functional degradation of the autophagosomal and normal autophagic flux. We also confirmed the increase in LC3-II turnover quantitatively by flow cytometry (Figures S3A and S3B).

Next, we investigated whether the increase in LC3-II turnover induced in butyrate macrophages is localized to intracellular Salmonella. To test this hypothesis, we performed confocal imaging by staining LC3 in control and butyrate macrophages infected with GFP-Salmonella (Figure 2L). Viable intracellular bacteria expressed a strong GFP signal. The degradation process was characterized by subsequent LC3 coating and loss of the GFP signal. Whereas in control macrophages a large proportion of Salmonella expressed a bright GFP signal, we found that in butyrate macrophages bacteria expressed a significantly lower GFP bright signal and a higher percentage of Salmonella are associated with strong LC3 coating (Figure 2L, details of the microscopy approach are described in detail in Figure S3C).
Figure 3. Butyrate Exposure Promotes an Antimicrobial Macrophage Phenotype

(A) Single-cell RNA-seq analysis of control and butyrate treated macrophages (n = 5,836 cells) from two healthy donors. The t-SNE projection shows the five subpopulations of macrophages that were identified with a graph-based clustering algorithm (see Methods).

(B) Quantitation of the proportions of control and butyrate macrophages ascribed to each of the sub-populations. Dashed lines indicate the sample pairs (shared donor identity).

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Moreover, we investigated the role of ROS generated by NADPH oxidase activity, an anti-microbial effector mechanism that typifies LC3-associated phagocytosis. In macrophages, the activity of the NADPH oxidase 2 (NOX2) enzyme complex is required for efficient recruitment of LC3 to phagosomes restricting bacterial colonization, growth in the cytosol, and elimination of invading bacteria (Huang et al., 2009). We measured NADPH oxidase activity by luminol-chemiluminescence on macrophages stimulated with phorbol 12-myristate 13-acetate (PMA) (Figure S3D) or after Salmonella infection (Figure S3E). In both conditions, butyrate macrophages produced significantly more reactive oxygen species (ROS) as measured by a dihydrorhodamine (DHR) assay after PMA stimulation or Salmonella infection compared to controls (Figures S3F and S3G).

**Calprotectin Mediates the Enhanced Antimicrobial Function of Butyrate Macrophages**

The antimicrobial effects of butyrate depended on sustained exposure during macrophage differentiation. Since butyrate has a well-known role as an HDACi, we reasoned that its effect on macrophage differentiation may be a consequence of epigenetic changes and gene expression affecting the heterogeneity of the resulting macrophage population. We therefore employed single-cell RNA-seq analysis, a method that facilitates the identification and transcriptomic characterization of previously unknown cell subtypes and states (Villani et al., 2017). We performed droplet-based single-cell RNA-seq analysis of control and butyrate macrophages from two healthy donors, retaining 5,981 cells for detailed analysis. We first performed an alignment between control and butyrate macrophages using an approach based on canonical correlation analysis (CCA) (Butler and Satija, 2018). We were able to align 98.7% of the cells, confirming the existence of broadly similar cell-states in both the control and butyrate macrophage populations. We identified five major clusters of macrophages (Figure 3A) and characterized these subpopulations by identifying marker genes for each cluster that were conserved between control and butyrate macrophages (selected examples shown in Figure 3C, and Table S1). The different clusters of macrophages were associated with distinct gene-expression profiles (cluster 0 lysosomal function; cluster 1 regulation of actin cytoskeleton and phagocytosis; cluster 2 phagosomal functions and antigen presentation; cluster 3 cell-cycle-associated genes). Cells in cluster 2 were characterized by genes associated with antimicrobial responses (e.g., S100A8, S100A9) and lysosomal functions (LYZ). Butyrate consistently increased the numbers of macrophages associated with the lysosomal (cluster 0) and antimicrobial phenotype (cluster 2) (Figure 3B).

To examine the influence of butyrate, we identified genes that were differentially expressed between control and butyrate macrophages within each of the clusters of differentiated (i.e., non-cell-cycle-associated) macrophages (see Methods). We found that butyrate predominantly induced changes in gene expression across the entire population of differentiated macrophages (Figure 3D, Table S2). Genes upregulated by butyrate were enriched for gene-ontology (GO) categories including “killing of cells of other organism” and “defense response to bacteria” (Figure 3E, Figure S4, Table S3). Butyrate downregulated genes associated with “cellular response to interleukin-1” and the classic complement (i.e., C1QA, C1QB, C1QC) activation genes in cells of cluster 2.

Overall, the single-cell data revealed a butyrate-induced antimicrobial signature characterized by the expression of S100A8, S100A9, S10012, LYZ, and FCN1 particularly in the cluster 2 subset of differentiated macrophages (Figure 4A and S8). Because these data are based on only two individuals, we next validated the induction of S100A8, S100A9, and S100A12 (Figure 4B) mRNA expression in butyrate macrophages compared to control macrophages by qPCR at baseline. Butyrate-induced S100A8 and S100A9 expression is not only observed at baseline but also in LPS stimulated cells suggesting that a short term inflammatory stimulus does not override the butyrate induced antimicrobial function (Figures S5A and S5B).

Analysis of control and butyrate macrophage supernatants by ELISA confirmed the significant increase in calprotectin protein secretion by butyrate macrophages (Figure 4C). Similarly, intracellular staining of the S100A8 and S100A9 proteins by flow cytometry shows a striking upregulation of calprotectin in macrophages after 5 days of differentiation with butyrate (Figure 4D). Quantification of calprotectin protein in the lysate of control and butyrate macrophages at steady state by immunoblot also confirmed a significant increase in butyrate macrophages (Figures 4E and 4F).

Calprotectin is a well-studied antimicrobial protein that is capable of enhancing bacterial killing via different mechanisms such as sequestration of metal ions such as zinc (Zn\(^{2+}\)) and manganese (Mn\(^{2+}\)) (Hood and Skaar, 2012; Kehl-Fie et al., 2011). To confirm the functional role of S100A8/9 in the elimination of invasive intracellular bacteria, we differentiated control and butyrate macrophages in the presence of siRNAs against S100A8 and S100A9 before performing a gentamicin protection assay. The significant silencing of both S100A8 and S100A9 genes (Figures S5C and S5D) found in butyrate macrophages treated with S100A8/S100A9 siRNA was associated with a significant increase of bacterial load compared to untreated or scrambled siRNA treated control and butyrate macrophages (Figure 4G). Together these results indicate that the presence of butyrate during monocyte to macrophage differentiation drives a synergistic program of antimicrobial LC3-associated host defense and production of antimicrobial peptides, which together promote cellular antimicrobial activity.
Butyrate Promotes Antimicrobial Activity in Macrophages through Its HDAC Inhibitory Function

Since butyrate can signal via G protein-coupled receptors (GPCRs) (Blad et al., 2012), we studied the role of GPCRs in butyrate-induced bacterial clearance. We found that mRNA expression of the known butyrate receptors FFAR2, FFAR3, and HACR2 genes were more highly expressed in myeloid populations than lymphoid and dendritic cell populations (Figures 5A–5C). However, macrophages differentiated in the presence of both butyrate and the GPCR inhibitor pertussis toxin (PT) still exhibited enhanced anti-bacterial activity (Figure 5D) suggesting that GPCRs are not required for butyrate induced antimicrobial functions.

We next tested the role of the histone deacetylase (HDAC) inhibitory function of butyrate in enhancing antimicrobial function. HDACs remove acetyl groups on specific lysine residues from histones and non-histone proteins regulating gene expression by modulating chromatin structure. We tested the ability of butyrate to inhibit HDAC activity in macrophages, using histone H3 and H4 acetylation as an indirect readout. Butyrate macrophages displayed a higher amount of acetylated H3 and H4 compared to controls (Figure 5E). Furthermore, butyrate macrophages showed a significant decrease of the tri-methylation of lysine 27 on histone 3 (3MeH3K27), associated with chromatin repression (Figure 5F), as well as increased acetylation of lysine 27 on histone 3 (AcH3K27), which is associated with more open chromatin (Figure 5G).

In support of these findings, treatment of macrophages with pan-HDAC inhibitors such as valproate, phenyl-butyrate, and vorinostat (SAHA) increased their antimicrobial activity (Figure 5H and 5J) and increased expression of S100A8 mRNA (Figure 5I and 5K).

We next investigated the specificity of butyrate-mediated HDAC inhibition in macrophages. HDACs are divided into 2 main classes: class I and class II (subdivided into IIa and IIb) (Haberland et al., 2009). We initially treated macrophages with TMP195, an inhibitor of HDAC class IIa (Davie, 2003) that has been used to activate tumor-associated macrophages (Guerriero et al., 2017) and with tubacin that targets HDAC class IIb. However, neither macrophages differentiated in the presence of TMP195 nor tubacin showed changes in bactericidal function (Figure 5L) or upregulation of S100A8 mRNA (Figure 5M).
Figure 5. Butyrate Promotes Antimicrobial Activity in Macrophages via HDAC Inhibitory Function

(A–C) Gene expression of hcar2 (A), ffmr2 (B), and ffmr3 (C) on flow cytometry sorted CD19+ B, CD8+ T, naive CD4+ T, memory CD4+ T, CD56+ CD3− NK cells, CD14+ monocytes, CD141+ DCs, and CD1c+ DCs from the blood of healthy donors. Each dot represents one donor.

(D) Gentamicin protection assay on control and butyrate macrophages, or butyrate macrophages differentiated in the presence of butyrate with pertussis toxin (PT).

(E) Protein expression of acetylated histone 3 (Ac-H3) and acetylated histone 4 (Ac-H4) in control and butyrate macrophages (data from 2 individual donors per condition).

(F and G) Intra-cellular expression of tri-methylated lysine27 on histone 3 (3MeH3K27) (F) and acetylated lysine 27 on histone 3 (AcH3K27) (G) protein on control and butyrate macrophages by flow cytometry.

(H, J, L, and N) Gentamicin protection assay on control macrophages, butyrate macrophages, and macrophages differentiated in the presence of valproate, phenylbutyrate (H), SAHA (J), TMP195, and tubacin (L), SBHA, 1-naphthoxyacetic acid (NA), or RGFP966 (N).

(I, K, M, and O) Gene expression of S100A8 in control macrophages, butyrate macrophages, and macrophages differentiated in the presence of valproate, phenylbutyrate (I), SAHA (K), TMP195, and tubacin (M), SBHA, 1-naphthoxyacetic acid (NA), or RGFP966 (O).

(P) Gentamicin protection assay in control and butyrate macrophages with and without HDAC3 siRNA-mediated gene silencing.

(Q) Extracellular acidification rate in control and butyrate macrophages, as well as in RGFP966 and TMP195 treated macrophages. Data represent the mean of three biological replicates.
suggesting that butyrate does not act via inhibition of class II HDAC. Next, we treated macrophages with several inhibitors that target class I HDACs: valproate (class I HDACi and reduces protein expression of HDAC2), SBHA (targeting HDAC1 and 3), 1-naphthoylhydroxamic Acid (NA: targeting HDAC8, 1, and 6), and RGFP966 (a HDAC3-specific inhibitor) (Jia et al., 2016). We found that macrophages differentiated with valproate and SBHA but not NA showed a significant reduction of CFU compared to control macrophages (Figure 5N). Similarly, the HDAC3 inhibitor RGFP966 induced elevated bacterial clearance (Figure 5N) and increased S100A8 mRNA expression (Figure 5O). Silencing of HDAC3 with siRNA resulted in significantly increased Salmonella clearance at baseline in control macrophages. Importantly, butyrate failed to enhance bacterial killing in macrophages with HDAC3 silencing (Figure 5P and Figure S5E). Together these results show that inhibition of HDAC3 is sufficient to induce the differentiation of macrophages with bactericidal functions and that butyrate’s anti-microbial effects on macrophages are dependent on HDAC3. Macrophages differentiated with the HDAC3 inhibitor RGFP966 but not TMP195 also showed reduced glycolysis similar to butyrate macrophages indicating inhibition of HDAC3 is upstream of both metabolic changes and anti-microbial responses in macrophages (Figure 5Q).

**Butyrate Treatment Promotes Antibacterial Activity in Intestinal Macrophages and Restricts Bacterial Translocation In Vivo**

To test whether the effects of butyrate on macrophage differentiation are relevant in vivo, we treated C57BL/6 mice with a daily oral dose of butyrate or water for 7 days. At day 7, we sorted colonic macrophages by flow cytometry and performed an ex vivo gentamicin protection assay. In line with our in vitro findings, colonic macrophages from butyrate-treated mice exhibited higher antimicrobial activity compared to controls (Figure 5G). Silencing of HDAC3 with siRNA resulted in significantly increased Salmonella clearance at baseline in control macrophages. Importantly, butyrate failed to enhance bacterial killing in macrophages with HDAC3 silencing (Figure 5P and Figure S5E). Together these results show that inhibition of HDAC3 is sufficient to induce the differentiation of macrophages with bactericidal functions and that butyrate’s anti-microbial effects on macrophages are dependent on HDAC3. Macrophages differentiated with the HDAC3 inhibitor RGFP966 but not TMP195 also showed reduced glycolysis similar to butyrate macrophages indicating inhibition of HDAC3 is upstream of both metabolic changes and anti-microbial responses in macrophages (Figure 5Q).

DISCUSSION

The intestinal immune system is highly adapted to provide host defense in the face of pathogens while retaining a mutualistic response with commensal bacterial. Bacterial fermentation products SCFA are major mediators of host-microbe cross talk in the intestine, controlling the development and maintenance of indigenous bacterial communities on the one hand and differentiation and maturation of intestinal tissue and immune cells on the other (Rooks and Garrett, 2016). Here we have identified a role for butyrate as a differentiation factor for monocyte-derived macrophages that enhances cell-intrinsic antimicrobial functions. We have shown that butyrate acted via its HDAC3 inhibition to alter metabolism and induce production of anti-microbial peptides leading to enhanced bactericidal function in vitro and in vivo.

Butyrate caused a profound metabolic and immunologic alteration in macrophages that is in many aspects opposite to the well-known pro-inflammatory LPS stimulation or “trained immunity” seen in pro-inflammatory M1 macrophages. LPS treated macrophages show increased glycolysis, inhibition of AMPK, increase of mTOR signaling, as well as reduced carbohydrate kinase-like protein (CARKL) mRNA that flux into the pentose phosphate pathway (Kelly and O’Neill, 2015). By contrast, macrophages differentiated in the presence of butyrate show reduced glycolysis, higher amounts of AMP and increased AMPK phosphorylation at Thr172, a residue that is critical for enzyme activity (Hardie et al., 2012). Butyrate macrophages also show reduced S6 phosphorylation, a surrogate marker for inhibition of mTOR. Reductions in mTOR, which is a known positive regulator of glycolytic enzymes such as hexokinase II, glyceraldehyde 3-phosphate dehydrogenase, and lactate dehydrogenase-B (Sun et al., 2011) may explain the reduced glycolysis observed in butyrate macrophages. In addition, butyrate treated macrophages show increased amounts of ribulose 5-phosphate and reduced intracellular glucose (despite normal glucose uptake) suggesting an

For the gentamicin protection assay, the percentage of CFU (of control) was calculated from mean value for control group. Each dot represents one independent donor. For pairwise comparison Mann-Whitney U test was performed and for multiple group comparisons a one-way ANOVA (Kruskal-Wallis test) was performed. *p < 0.05, **p < 0.01, and ***p < 0.001. Please see Figure S5E.
increased flux toward the pentose phosphate pathway. An increased flux toward the pentose phosphate pathway could contribute to the generation of NADPH, which may fuel the increased NADPH-oxidase-dependent ROS observed in butyrate macrophages both at baseline and also after Salmonella infection. Consistent with previous data that increased AMP kinase activity inhibits mTOR signaling and induces autophagy (Kelly and O’Neill, 2015; Shimobayashi and Hall, 2016), we found increased LC3 and increased bacterial clearance as a marker of effective autophagy in butyrate differentiated macrophages (Kim and Guan, 2015). Our results suggest an LC3-associated process since butyrate macrophages express high amounts of lipidated LC3-II protein detected by immunoblot, flow cytometry, and microscopy. However, we cannot differentiate whether the effects of butyrate are linked to canonical autophagy or non-canonical LC3-associated phagocytosis pathways (Martinez et al., 2015).

Single-cell RNA-sequencing identified five clusters of gene expression among human monocyte-derived macrophages with enrichment of distinct cellular processes reiterating the emerging concept of phenotypic and functional heterogeneity. Addition of butyrate increased expression of a number of anti-bacterial and...
host defense genes supporting the marked increase in bacterial killing in butyrate macrophages at the population level. However, a butyrate-induced antimicrobial signature involving the expression of S100A8, S100A9, S10012, LYZ, and FCN1 was particularly pronounced in a subset of differentiated macrophages (cluster 2). The modulation of metabolic regulation toward LC3-associated processes in conjunction with induction of multiple antimicrobial genes within those cells provides the machinery for intracellular bacterial segregation and effector mechanisms for killing. Calprotectin (a heterodimer of S100A8 and S100A9) mRNA and protein was detected in butyrate macrophages. Gene silencing of calprotectin alleviated the increased anti-bacterial properties of butyrate macrophages, illustrating the functional importance of this antimicrobial peptide. Calprotectin can inhibit the growth of intracellular pathogens by a number of mechanisms including chelation of bivalent cations (Zaia et al., 2009), activation of NADPH oxidase (Kerkhoff et al., 2005), or induction of lipidated LC3-II (Wang et al., 2015).

We have shown that HDAC3 inhibition by butyrate (or the specific HDAC3i RGFP966) drove the differentiation of macrophages, altered their metabolism, and enhanced gene expression of antimicrobial peptides uncoupled from increased inflammatory cytokine production. Butyrate has been shown to act as an HDACi to inhibit acute LPS stimulated inflammatory cytokine production by murine macrophages in vitro (Chang et al., 2014). In that setting HDAC3 deficiency or inhibition has anti-inflammatory function (Chen et al., 2012; Leus et al., 2016). Lyz2−cre+/Hdac3−/+ macrophages also polarize more strongly toward an M2 phenotype when stimulated with IL-4 (Mullican et al., 2011). However, that function is context dependent and requires IL-4 signaling. Our findings complement and extend previous studies by showing that butyrate through its HDAC3i function can also induce the differentiation of a specialized macrophage anti-microbial state in the absence of changes in inflammatory cytokine production. Together these studies support a model in which butyrate imprints a non-inflammatory and antimicrobial program in macrophages that promotes intestinal homeostasis. This is consistent with the concept that butyrate-producing commensal bacteria shape host microbial crosstalk to promote a stable relation-ship, avoiding the disruptive effects of inflammation on the ecosystem. This is an important mechanism, since genetic defects in bacterial handling and reduction in butyrate producing bacteria have been linked to IBD (Baxt and Xavier, 2015; Cadwell, 2016; Frank et al., 2007). This also fits with the observation that human intestinal macrophages showed reduced inflammatory potential compared to peripheral blood monocytes (Bujko et al., 2018). Our results show that mice pre-treated with butyrate exhibit a reduction of systemic dissemination when orally infected with C. rodentium or Salmonella. Although we cannot exclude that in the in vivo setting butyrate promotes antimicrobial barrier function via several mechanisms, macrophages sorted from the colon of butyrate-fed mice display an increase in bactericidal function and an upregulation of S100A8 mRNA suggesting that these cells contribute to antimicrobial homeostasis.

Epidemiological studies suggest that early life exposure to antibiotics is a risk factor and a high-fiber diet (associated with increased luminal butyrate concentrations) is a protective factor in IBD (Ananthakrishnan et al., 2018; Cushing et al., 2015). In a model system, broad spectrum antibiotics causes butyrate depletion and butyrate-responsive macrophage and T cell dysfunction (Scott et al., 2018). Restoring antimicrobial function via butyrate in intestinal macrophages may therefore be a universal mechanism to prevent or treat IBD. It is feasible to pharmacologically increase butyrate concentrations via enema in vivo in portal vein blood to 92.2 μmol/L (van der Beek et al., 2015). Although it is not yet clear whether enemas containing butyrate or a cocktail of SCFAs can ameliorate intestinal inflammation (Cushing et al., 2015), treatments that harness butyrate effector mechanisms might have the potential to prevent IBD or reduce relapse activity. Our findings suggest clear differences between the anti-tumor inflammatory macrophages induced by the class II HDACi TMP195 and antimicrobial macrophages induced by HDAC3 targeting inhibitors suggesting a concept of differential imprinting of macrophage function via selective HDAC inhibition. It remains to be determined whether selective HDAC3 inhibitors would have an advantage beyond their natural counterpart butyrate. Indeed, HDAC3 inhibition might induce complex effects on other cell types such as those present in the epithelium (Alenghat et al., 2013).

In conclusion, our results demonstrate that butyrate directs the differentiation of homeostatic macrophages that possess strong antimicrobial activity and play an important role in preventing the dissemination of bacteria beyond the intestinal barrier. Butyrate educates developing macrophages via HDAC3 inhibition by regulating their metabolic and transcriptional program. This has implications for prevention and therapy of disorders that are associated with intestinal inflammation, as well as systemic infection.

**STAR METHODS**

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SUPPLEMENTAL INFORMATION
Supplemental Information includes six figures and three tables and can be found with this article online at https://doi.org/10.1016/j.immuni.2018.12.018.

ACKNOWLEDGMENTS
We thank all volunteers and blood donors for participation in this study. We thank Sebastian Rogati for technical assistance, Helen Ferry and Jonathan Webber for flow cytometry cell sorting, and the staff of our animal houses. We acknowledge the contribution of the BRC Gastrointestinal biobank (11/YH/0020, 16/YH/0247), supported by the National Institute for Health Research (NIHR) Oxford Biomedical Research Centre (BRC). This work was supported by the Welcome Trust UK (095688/Z/11/Z: F.P., D.G.W.J., and D.R.S.). The BBSRC (BB/I005609/1: F.P., C.V.A.-C, J.S., M.C.), the Leona M. and Harry H. Helmsley Charitable Trust (F.P., H.H.U, J.S, M.C.), the Kennedy Trust for Lupus Research (K.C.R.-A., N.I., S.N.S., and J.M.), the Crohn’s & Colitis Foundation of America (H.H.U, F.P, J.S.), and the BRC Gastrointestinal biobank. We acknowledge the contribution of the BRC Gastrointestinal biobank (11/YH/0020, 16/YH/0247), supported by the National Institute for Health Research (NIHR) Oxford Biomedical Research Centre (BRC). This work was supported by the Wellcome Trust UK (095688/Z/11/Z: F.P., D.G.W.J., and D.R.S.).

DATA AND SOFTWARE AVAILABILITY
Supplemental Information includes six figures and three tables and can be found with this article online at https://doi.org/10.1016/j.immuni.2018.12.018.

AUTHOR CONTRIBUTIONS
J.S., S.P., I.A., M.C., F.F., A.C., J.M., D.G.W.J. performed experiments and K.C.R.-A., N.I., S.N.S. and J.M. bioinformatics analyses. S.N.S. supervised the single-cell analysis; C.V.A.-C. supervised initial experiments; F.P. and H.U. supervised the study. All authors discussed the data and manuscript.

DECLARATION OF INTEREST
None of the authors has a conflict of interest related to this article. H.H.U. received research support or consultancy fees from UCB Pharma, Eli Lilly, Boehringer Ingelheim, Pfizer, Celgene, and AbbVie. F.P. has received research support or consultancy fees from GSK, UCB Pharma, Medimmune, Janssen, Roche, and Eli Lilly.

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## STAR METHODS

### KEY RESOURCES TABLE

| REAGENT or RESOURCE | SOURCE | IDENTIFIER |
|---------------------|--------|------------|
| **Antibodies**      |        |            |
| Anti-CD45RO-APC Cy7/biotin (Clone UCHL1) | BioLegened | Catalog# 304228; RRID:AB_10895897 |
| CD45RA PE Cy7/biotin (Clone HI100) | BioLegened | Catalog# 304125; RRID:AB_10709440 |
| CD8-V500/biotin (Clone RPA-T8) | Biosciences | Catalog# 560775; RRID:AB_1937333 |
| CD19- BV650/biotin (Clone HIB19) | BioLegend | Catalog# 302238; RRID:AB_2562097 |
| TAAAD | BioLegend | 420403 |
| Annexin V-BV510 | BioLegend | 640937 |
| Anti-GLUT1 (Clone FAB1418P) | RnD Systems | Catalog# FAB1418P; RRID:AB_2191040 |
| Anti-CD14 (Clone M5E2) | BioLegend | Catalog# 301804; RRID:AB_314186 |
| Anti-CD11c (Clone B-Ly6) | BD Biosciences | Catalog# 657713; RRID:AB_2760137 |
| Anti-HLA-DR (Clone G46-6) | BD Biosciences | Catalog# 561224; RRID:AB_10563765 |
| Anti-Foxp3-PE-eFluor610 (Clone FJK-16s) | eBioscience | Catalog# 61-5773-80; RRID:AB_2574623 |
| Anti-CD4-BV785 (Clone RM4-5) | BioLegend | Catalog# 100552; RRID:AB_2563053 |
| Anti-CD45-BV650 (Clone 30-F11) | BioLegend | Catalog# 103151; RRID:AB_2565884 |
| Anti-CD3-PECy7 (Clone 145-2C11) | BioLegend | Catalog# 100320; RRID:AB_312685 |
| fixable viability dye | eBioscience | Catalog# 65-0865-14 |
| Anti-S100A8 (Clone CF-145) | eBioscience | Catalog# 50-9745-42; RRID:AB_2574354 |
| Anti-S100A9 (Clone MRP-14) | BioLegend | Catalog# 350706; RRID:AB_2564008 |
| Anti-phosphor-S6 (ser235, ser236) (Clone cupk43k) | eBioscience | Catalog# 12-9007-42; RRID:AB_2572667 |
| Anti-acetyl-histone H3 (lys27) (Clone D5E4) | New England Biolabs | Catalog# 8173; RRID:AB_10949503 |
| Anti-tri-methyl-histone H3 (lys27) (Clone C36B11) | New England Biolabs | Catalog# 9733; RRID:AB_2616029 |
| Anti-LC3B (Clone D11) | Cell Signaling | Catalog# 3868; RRID:AB_2137707 |
| Anti-β-actin (Clone 13E5) | Cell Signaling | Catalog# 5125; RRID:AB_1903890 |
| Anti-phosphor-S6 ribosomal protein (Ser235/236) (Clone D57.2.2E) | Cell Signaling | Catalog# 4858; RRID:AB_916156 |
| Anti-S100A8 (Clone EPR3554) | Abcam | Catalog# ab92331; RRID:AB_2050283 |
| Anti-acetylated-H3 (Clone ab47915) | Abcam | Catalog# ab47915; RRID:AB_873860 |
| Anti-acetylated-H4 (Clone EPR16606) | Abcam | Catalog# ab177790; RRID:AB_2732882 |
| Anti-P62 lck (Clone 2/P62 LCK Ligand) | BD Bioscience | Catalog# 610833; RRID:AB_398152 |
| HRP-conjugated secondary antibody (anti-rabbit) | Cell Signaling | Catalog# 7074S |
| HRP-conjugated secondary antibody (anti-rabbit) | Cell Signaling | Catalog# 7076S |
| **Bacterial strains** |        |            |
| Salmonella enterica serovar Typhimurium | D. Holden (Imperial College, University of London, UK) | NCTC 12023 |
| CD-associated adherent invasive Escherichia coli (AIEC) | Arlette Darfeuille-Michaud lab France | LF82.30 |
| Staphylococcus aureus | National collection of type cultures | NCTC 6571 |
| Citrobacter rodentium | Gad Frankel in Imperial College London | ICC169 |
| Salmonella (mouse) | Gordon Dougan lab, Cambridge university | SL1344 |
| **Chemicals** |        |            |
| Human M-CSF | Preprotech |          |
| Sodium Butyrate | Sigma-Aldrich | 303410-100G |
| Sodium Acetate | Sigma-Aldrich | S2889-250G |
| Sodium Propionate | Sigma-Aldrich | P1880 |

(Continued on next page)
Human monocytes were isolated from leukocyte cones of healthy blood donors. Peripheral blood mononuclear cells (PBMC) were obtained by ficoll gradient. Monocyte-derived macrophages were generated using adherence method selection and M-CSF differentiation. Whole PBMC (50×10^6) were plated in RPMI-1640 medium for 90 min. After 2 washes with PBS, adherent monocytes were differentiated into macrophages over a 5 day period in presence of 100 ng/mL M-CSF in RPMI supplemented with 10% fetal calf serum (FCS) (Sigma-Aldrich), 100 U/mL penicillin, 100 μg/mL streptomycin, 30 mM HEPES, and 0.05 mM β-mercaptoethanol.

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**HUMAN MONOCYTES, DIFFERENTIATION AND CELL CULTURE**

Human monocytes were isolated from leukocyte cones of healthy blood donors. Peripheral blood mononuclear cells (PBMC) were obtained by ficoll gradient. Monocyte-derived macrophages were generated using adherence method selection and M-CSF differentiation. Whole PBMC (50x10^6) were plated in RPMI-1640 medium for 90 min. After 2 washes with PBS, adherent monocytes were differentiated into macrophages over a 5 day period in presence of 100 ng/mL M-CSF in RPMI supplemented with 10% fetal calf serum (FCS) (Sigma-Aldrich), 100 U/mL penicillin, 100 μg/mL streptomycin, 30 mM HEPES, and 0.05 mM β-mercaptoethanol.
REAGENTS

All chemicals Butyrate (1mM), Acetate (1mM), Propionate (1mM), SAHA (1μM), SBHA (20μM), TMP195 (10 μM), RGFP966 (20 μM), 1-NA (10 μM), tubacin (1μM), 1-naphthohydroxamic acid, bafilomycin A1, 3-MA, MHY1485) were purchased from Sigma-Aldrich, unless specified otherwise. TMP195 was acquired from Cellagen technology and Cayman chemical respectively. Recombinant human and murine M-CSF were purchased from PeproTech.

BACTERIAL STRAINS AND GENTAMICIN PROTECTION ASSAY

Gentamicin protection assay was performed with the following strains: Salmonella enterica serovar Typhimurium (Salmonella)-expressing green-fluorescent protein (GFP) (NCTC 12023), CD-associated adherent invasive Escherichia coli (AIEC) reference strain LF82.30, Staphylococcus aureus (NCTC 6571). All strains were used at an MOI of 10 unless specified otherwise. For the gentamicin protection assay, macrophages were infected for 1h with Salmonella or AIEC or S. aureus or C. rodentium followed by gentamicin treatment for 2h. Cells were then lysed in 1% triton buffer and the lysate was plated on agar plates. Results are presented as absolute CFU count or % of mean control.

SEAHORSE ASSAY

Extra cellular acidification rate (ECAR) of the control and butyrate treated Macrophages was quantified by using a XF 96 extracellular flux analyzer (Seahorse Bioscience). 100,000 macrophages / well was plated in Seahorse base media was supplemented with 1% FCS, 1mM glutamine and 2mM sodium pyruvate. Plate were incubated in a Co2 free incubator at 37°C and transferred to Seahorse machine for ECAR quantification. The assay was performed on 8 donors (biological replicates) with 5-8 technical replicates per donor. Similarly, for the mito-stress test, oxygen consumption rate (OCR) was quantified using a XF 96 extracellular flux analyzer (Seahorse Bioscience) as per the manufactures protocol. Base media for mito-stress was supplemented with 1% FCS, 1mM glutamine, 2mM sodium pyruvate and 10mM glucose.

METABOLOMICS ANALYSIS BY LC-MS

Metabolite extraction from cells

Metabolites were extracted from approximately 1x10⁶ cells (grown in cell culture dishes) by addition of 500 μL of ice cold 80% aqueous methanol. The supernatants were combined and filtered using a 3 kD ultrafilter (Millipore), dried in a SpeedVac and subsequently stored at −80°C. On the day of analysis, the dried extracts were re-constituted in 60 μL of ice cold 80% aqueous methanol. A quality control (QC) sample was made by combining 5 μL of each sample. This was injected at the start of the sequence and subsequently every 10 samples throughout the LC-MS/MS analyses.

LC-MS/MS analysis

Each sample was analyzed using two different LC-MS/MS methods utilizing two separate chromatographic systems. The first method used ion-chromatography coupled directly to Q-exactive HF Hybrid Quadrupole-Orbitrap mass spectrometer (IC-MS) (Thermo Scientific San Jose, CA). The second method utilized reversed-phase ultra-high performance chromatography (UHPLC) coupled directly to the same Q-exactive HF Hybrid Quadrupole-Orbitrap mass spectrometer (Thermo Ultimate 3000, Thermo Scientific, San Jose, CA). Both methods have been published previously and further details can be found in (French et al., 2018; Riffelmacher et al., 2017).

Data processing

Raw data files were processed using ProgenesisQI (Waters, Elstree, UK). This involved alignment of retention times, peak picking by identification of the presence of natural abundance isotope peaks, characterizing multiple adduct forms and identification of metabolites using our in-house database of authentic standards. Retention times, accurate mass values, relative isotope abundances and fragmentation patterns were compared between authentic standards and the samples measured. Identities were accepted only when the following criteria were met: < 5ppm differences between measured and theoretical mass (based on chemical formula), < 30 s differences between authentic standard and analyte retention times, isotope peak abundance measurements for analytes were > 90% matched to the theoretical value generated from the chemical formula. Where measured, fragmentation patterns were matched to the base peak and two additional peak matches in the MS/MS spectrum to within 12ppm. The top 10 data directed fragmentation method was not always able to provide fragment ions for all ions measured in the MS 1 spectrum.

Data analysis

Principal Component Analysis (PCA) was performed using Progenesis QI. Fold change, % CV and p values were generated automatically in progenesis QI and verified manually using a normalized abundance output and Excel. Heatmaps were generated manually.
using the verified fold-change output. p values were generated using ANOVA (independent conditions). Statistical differences were annotated according to the scale *p < 0.05, **p < 0.01, ***p < 0.001.

Unsupervised clustering analysis was performed using the ClustVis web tool (https://biit.cs.ut.ee/clustvis/).

**AMPK ELISA**

Monocytes were differentiated in the presence of M-CSF and butyrate or left untreated for 5 days. For quantification of pAMPK phosphorylation 100,000 macrophages were plated in flat bottom plates and AMPK quantification was performed at baseline after cell lysis as per the manufactures protocol (KHO0651; Thermo scientific).

**ANNEXIN V AND 7AAD STAINING**

For the Annexin V and 7AAD staining, 250,000 cells were harvested after 5 days of differentiation. In the case of Salmonella infection, cells were infected at an MOI of 10 for 1 h prior to staining. Annexin V and 7AAD staining was performed as per the manufactures protocol.

**MEASUREMENT OF REACTIVE OXYGEN SPECIES**

Production of reactive oxygen species by macrophages was evaluated with the chemiluminescence probe L-012 (100 µM, Wako laboratories, Japan) in opaque white 96-well plates. Cells were activated with 100ng/mL of PMA or infected with Salmonella (MOI 10) for 30 min. The luminescence was recorded every 2 min for 90 min with a plate reader (FLUOstar OPTIMA, BMG labtech). For the flow cytometric dihydrorhodamine (DHR) assay, macrophages were treated with DHR (2.5 µg/mL) and stimulated with/without 100 ng/mL PMA and analyzed by flow cytometry.

**CONFOCAL MICROSCOPY**

Quantification of GFP positive bacteria and Salmonella-associated LC3 was performed by confocal microscopy. Control and butyrate treated macrophages (1x10⁵) were seeded in 8-well chamber slides (Sarstedt) and were infected with GFP Salmonella Typhimurium at an MOI of 20 h. Cells were then treated for an additional 1 h with gentamicin (100µg/ml) to kill extracellular Salmonella. Cells were fixed with 2% paraformaldehyde and then permeabilized with 0.1% Triton X-100 for 10 min. Fixed and permeabilized cells were stained with anti-LC3 (clone PM036, MBL) and secondary antibodies (Alexa Fluor 568 goat anti-rabbit IgG, Life technologies). Cellular and bacterial DNA was stained with DAPI. Finally, images of macrophages were acquired as a stack of images collected at 0.5 µm intervals at 63x magnification with a Zeiss 510 or 780 inverted confocal microscope (ZEN2009 or ZEN2011 software). A minimum of 100 infected cells were evaluated and quantified with ImageJ software. For the microscopy quantification of Salmonella degradation, bacteria were classified into GFPbright LC3negative/dim (i.e., early stage Salmonella infection with intact GFP signal) and GFPdimLC3bright  (LC3 coated Salmonella with quenched or degraded GFP signal). In addition there are intermediate stages of GFPdimLC3dim (Intermediate phase) and GFPnegativeLC3negative Salmonella (end stage degradation with bacterial DNA remnant). DAPI stain was used to identify intracellular bacteria. The microscopy acquisition setting was used to identify GFPbright and LC3bright bacteria whereas the enhanced brightness setting of each image was used to confirm the bacterial DNA content and to identify intermediate stages.

**FLOW CYTOMETRY**

Phenotyping and characterization of activation markers of human monocytes-derived macrophages were performed by flow cytometry. Cells were harvested, washed and counted before being incubated for 20 min at 4°C in PBS with 5% BSA containing the following surface antibodies: CD14 (clone M5E2) (Biolegend), CD11c (clone B-Ly6), HLA-DR (G46-6) (BD Biosciences) and fixable viability dye (eBioscience). Following surface staining, cells were fixed and permeabilized with the Cytofix/Cytoperm Fixation/Permeabilization Solution Kit (BD Biosciences) according to the manufacturer’s instructions. Cells were intra-cellularly stained with S100A8 (clone CF-145, eBioscience), S100A9 (clone MRP1H9, Biolegend), phospho-S6 (ser235, ser236) (clone cupk43k, eBioscience), acetyl-histone H3 (lys27) (clone D5E4) (New England biolabs) and tri-methyl-histone H3 (lys27) (clone C36B11) (New England biolabs). LC3-turnover was assessed by flow cytometry with the Autophagy Detection Reagent according to manufacturer’s instructions (Merck Millipore). GLUT1 receptor expression was quantified by flow cytometry (n = 4 healthy donors). 250,000 macrophages were surface stained in the flow cytometry staining buffer for 20 min on ice. Glucose uptake was quantified using 2-NBDG, a fluorescent glucose analog. Cells were incubated with 10µM 2-NBDG in RPMI + 10% FCS media at 37°C. All cells were acquired on a LSRII or a Fortessa (BD Biosciences). All analysis was performed using FlowJo software (Tree Star, Ashland, OR).
CD45+CD3+CD4+FOXP3+ T CELLS QUANTIFICATION

Cells were isolated as described in Methods (Isolation of mouse colonic macrophages). Staining was performed on uninfected mice or 48 h after *Salmonella* infection. Butyrate-treated mice received butyrate 5 days before and during infection in the drinking water. Lamina propria and spleen CD45+CD3+CD4+FOXP3+ T cells were analysed by flow cytometry.

ANALYSIS OF SINGLE-CELL TRANSCRIPTOMICS DATA

Single-cell RNA-sequencing libraries were generated using the 10x Genomics Single Cell 3’ Solution (version 2) and subjected to Illumina sequencing (HiSeq 4000). The computational workflow used to analyze the 10x Genomics data is available at https://github.com/sansomlab/tenx. Briefly, reads were aligned using 10x Genomics Cell Ranger pipeline (version 2.0.2) and human reference sequences (version 1.2.0). To circumvent known index-hopping issues with the HiSeq 4000 platform (Sinha et al., 2017), cell barcodes common to more than one sample were removed from the aggregated count matrix. UMI counts were randomly down-sampled so as to normalize the median number of per-cell counts between the samples. Data were then processed using the Seurat CRAN package (version 2.2.0). Cells with >20% mitochondrial reads or fewer than 500 genes were excluded from the analysis. Per-cell counts were normalized, scaled and the effects of total UMI count, percentage of mitochondrial UMI count, and donor (within the control and butyrate conditions) regressed out. Cells from the two conditions were then aligned using an approach based on canonical correlation analysis (Butler & Satija, 2018). We retained the first 13 CCA components, and discarded cells for which the variance explained by CCA was < 2-fold (i.e., ratio < 0.5) of that observed with PCA. Clusters of cells were identified using the FindClusters function of the Seurat package (original Louvain algorithm, resolution = 0.3) and visualized by t-Distributed Stochastic Neighbor Embedding (t-SNE) projection (perplexity = 20, Figure 3A) (Van der Maaten and Hinton, 2008). For each cluster of cells, markers were identified using the MAST test (Finak et al., 2015) as implemented in the Seurat package (default parameters; Table S1). Conserved cluster markers (Figure 3C) were identified as those achieving a maximum BH adjusted P value of < 0.05 when tested within each of the samples separately. Gene set enrichment analysis (Figure 3E) was performed using Fisher’s Exact Test (FET), Biological Process gene sets obtained from Gene Ontology (GO) databases (Ashburner et al., 2000; Kanehisa and Goto, 2000; Liberzon et al., 2011; Subramanian et al., 2005), and a gene universe that comprised of genes expressed in the differentiated Macrophages (n = 11,218). To identify genes differentially expressed between control and butyrate macrophages, we first summed the counts for each samples’ cells within each of the clusters. The DESeq2 package (Love et al., 2014) was then used to model the replicated, paired design and to perform a test for differential gene expression between control and butyrate macrophages within each of the clusters. The accession number for the single cell sequencing reported in this paper is Gene Expression Omnibus: GSE111049.

QUANTITATIVE PCR

RNA was isolated from macrophages, or from flow cytometry-sorted cells using the RNeasy Mini kit (QIAGEN, Manchester, UK) according to the manufacturer’s instructions. cDNA synthesis was performed using High-Capacity cDNA Reverse Transcriptase (Applied Biosystems, Life Technologies, Paisley, UK). qPCR were performed using TaqMan gene expression assays (Life Technologies) and TaqMan Universal PCR master mix. qPCR were run on the CFX96 detection system (Bio-Rad Laboratories, Hemel Hempstead, UK) and gene expression for each sample were normalized to RPLPO for human reference gene or HPRT for mouse reference gene and the differences were determined using the 2^ΔΔC(t) calculation.

QUANTITATION OF CYTOKINE SECRETION

Calprotectin was measured by ELISA (Biolegend) in the culture supernatants of control and butyrate macrophages after 5 days of differentiation by ELISA accordingly to the manufacturer’s instruction and normalized to the cell count.

TRANSFECTION OF PRIMARY HUMAN MACROPHAGES

The Accell SMARTpool siRNA were used to target human S100A8, S100A9 and HDAC3 for siRNA-mediated gene silencing in macrophages according to the manufacturer’s instructions (Dharmacon).

IMMUNOBLOT

Cells were lysed and the protein extract was quantified by BCA (Thermo Scientific). Equal amounts of total cell lysates (20 μg) was run on SDS-PAGE gels (4%–20%) (Bio-Rad), followed by transfer to polyvinylidene difluoride membranes. Proteins were detected using primary antibodies against LC3B (D11, Cell signaling), β-actin (13E5, Cell signaling), phospho-S6 ribosomal protein (Ser235 and/or 236) (D57.2.2E, Cell signaling), S100A8 (EPR3554, Abcam), acetylated-H3 (ab47915, Abcam) and acetylated-H4 (EPR16606, Abcam), P62 lck (clone 3/P62, BD bioscience) and detected using HRP-conjugated secondary antibodies.
ISOLATION OF LYMPHOID AND MYELOID POPULATIONS FROM HUMAN BLOOD

PBMC from healthy donors were isolated as detailed above and stained for surface markers in PBS with 0.1% bovine serum albumin buffer allowing MACS enrichment and FACS sorting. CD4+ T cells were enriched by depleting CD8+, CD19+, CD56+ and CD14+ cells by MACS and then stained using anti-CD4, anti-CD45RA and anti-CD45RO to sort naive CD4+CD45RA+ and memory CD4+CD45RO+ T cells by FACS. B cells, CD8+ T cells and NK cells were sorted by FACS using anti-CD19 for B cells, anti-CD8 for CD8+ T cells or anti-CD56+ for NK cells. CD141high and CD1c+ DC populations were enriched by depleting CD3+, CD19+, CD56+ and CD14+ cells by MACS and FACS sorted to isolate HLA-DRhigh CD11chigh CD141high for CD141+ DC or HLA-DRhigh CD11chigh CD141+ CD1c+ for CD1c+DC. All cells were sorted using an AriaIII flow cytometer (BD Biosciences) to a purity of > 95%. CD14+ monocytes were obtained by anti-CD14 bead enrichment using MACS sort.

EXPERIMENTAL MODEL AND SUBJECT DETAILS

Animals were bred and maintained under specific pathogen-free conditions. Animal experiments were performed in accordance with the UK Scientific Procedures Act (1986) under a Project License (PPL) authorised by the UK Home Office. 8- to 12-week-old inbred female C57BL/6J were used for experiments and were bred and maintained at the University of Oxford. Mice were routinely screened for the absence of pathogens, and were kept in individually ventilated cages with environmental enrichment. For experiments involving infection and/or butyrate administration, all mice in individual cages received the same treatment. To reduce cage effects each treatment was given to more than one cage in each experiment.

SODIUM BUTYRATE TREATMENT AND INFECTION

Mice received either saline, as a control, or sodium butyrate (150mM final concentration) (Sigma-Aldrich, St. Louis, MO) in their drinking water for 7 days before the sort of colonic macrophages at steady state. In case of Salmonella Typhimurium 1344 def AroA and Citrobacter rodentium infection, mice received saline as a control or sodium butyrate in drinking water for 5 days and 3 days respectively before infection. For the infection, mice were intragastrically gavaged with nalidixic acid resistant Citrobacter rodentium (ICC169) (1x10⁹) or Salmonella Typhimurium (1x10⁹). Mice weight was monitored throughout the duration of experiment (7 days for Citrobacter rodentium and 2 days for Salmonella Typhimurium).

HISTOLOGY SCORE

Samples of the proximal colon were collected and fixed in buffered 10% of 36% formalin solution to assess the severity of colitis. Haematoxylin and eosin (H&E) staining was performed on 4–5 µm paraffin-embedded sections and inflammation was assessed using previously published criteria (Izcue et al., 2008). Each sample was graded semiquantitatively based on the following features: cellular infiltration of the lamina propria, epithelial hyperplasia and goblet cell depletion, percentage of the section affected, and markers of severe inflammation (submucosal inflammation, crypt abscesses) from 0 to 4 in each of the 4 parameters. Samples were scored by two individuals blinded and scores for each criterion were added to give an overall score for each sample of 0–12.

ISOLATION OF MOUSE COLONIC MACROPHAGES

Colons from 10 mice gavaged with PBS and with sodium butyrate were prepared as described (Arnold et al., 2016). Briefly colons were opened, luminal content removed and washed, and tissue cut into pieces. Pieces were washed twice in RPMI-1640 medium supplemented with 5% fetal bovine serum and 5mM EDTA at 37°C with shaking to remove epithelial cells. Tissue was then digested in RPMI medium containing 1mg/mL type VIII collagenase (Sigma-Aldrich, Gillingham, UK), 0.5mg/mL DNase I and 5% fetal bovine serum. Cells were then layered on a 40/80% Percoll gradient, centrifuged, and the colonic leukocytes at the gradient interface were recovered and were stained in phosphate-buffered saline with 0.1% bovine serum albumin buffer with a combination of the following antibodies: CD45 (30-F11), CD11c (N418), MHCII (M5/114.15.2), F4/80 (BM8); CD103 (M290), CD11b (M1/70), CD64 (X54-5/7.1). Before the sort, DAPI was added to differentiate viable from dead cells. Macrophages were defined as (CD11b+ CD64+ F4/80+ MHCII+ CD103-) positive. All cells were sorted on an AriaIII (BD Biosciences) flow cytometer to a purity of > 95%.

BONE MARROW DERIVED MACROPHAGES

In brief, bone marrow was extracted from the femur of mice, washed, cells were counted and seeded at the concentration of 7x10⁶ per 10cm dish in 10 mL of RPMI supplemented with 50 ng/mL murine M-CSF with 10% FCS (Sigma-Aldrich), 100 U/mL penicillin, 100 µg/mL streptomycin, 30 mM HEPES for 7 days with or without butyrate (1mM).
PHAGOCYTOSIS ASSAY

Macrophage phagocytic activity was measured by two different methods: Firstly by using *Escherichia coli* (K-12 strain) BioParticles™, Alexa Fluor™ 488 conjugate as per manufacture’s instruction; Secondly, macrophages were infected with GFP-*Salmonella* and incubated for indicated time (15 min, 30 min, 45 min, 60 min). Cells were washed and the fluorescence due to *Salmonella* attached to the outside of the cells was quenched with trypan blue solution (4%). The fluorescence was evaluated by flow cytometry.

STATISTICS

Depending on the dataset, a Mann–Whitney nonparametric-test or Kruskal Wallis test (one-way Anova analysis) was used to calculate significance between groups. Graph generation and statistical analyses were performed using Prism version 5.0d software (GraphPad, La Jolla, CA). Differences were considered statistically significant when p < 0.05.

DATA AND SOFTWARE AVAILABILITY

The accession number for the single cell sequencing reported in this paper is Gene Expression Omnibus: GSE111049. This accession number will be accessible to readers upon publication of the manuscript.
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Title: The Short Chain Fatty Acid Butyrate Imprints an Antimicrobial Program in Macrophages

Date: 2019-02-19

Citation: Schulthess, J; Pandey, S; Capitani, M; Rue-Albrecht, KC; Arnold, I; Franchini, F; Chomka, A; Ilott, NE; Johnston, DGW; Pires, E; McCullagh, J; Sansom, SN; Arancibia-Carcamo, C; Uhlig, HH; Powrie, F, The Short Chain Fatty Acid Butyrate Imprints an Antimicrobial Program in Macrophages, IMMUNITY, 2019, 50 (2), pp. 432 - +

Persistent Link: http://hdl.handle.net/11343/234091

File Description: Published version