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Diacylglycerol kinase synthesized by commensal *Lactobacillus reuteri* diminishes protein kinase C phosphorylation and histamine-mediated signaling in the mammalian intestinal epithelium

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*Lactobacillus reuteri* 6475 (Lr) of the human microbiome synthesizes histamine and can suppress inflammation via type 2 histamine receptor (H2R) activation in the mammalian intestine. Gut microbes such as Lr promote H2R signaling and may suppress H1R proinflammatory signaling pathways in parallel by unknown mechanisms. In this study, we identified a soluble bacterial enzyme known as diacylglycerol kinase (Dgk) from Lr that is secreted into the extracellular milieu and presumably into the intestinal lumen. DgkK diminishes diacylglycerol (DAG) quantities in mammalian cells by promoting its metabolic conversion and causing reduced protein kinase C phosphorylation (pPKC) as a net effect in mammalian cells. We demonstrated that histamine synthesized by gut microbes (Lr) activates both mammalian H1R and H2R, but Lr-derived Dgk suppresses the H1R signaling pathway. Phospho-PKC and IκBα were diminished within the intestinal epithelium of mice and humans treated with wild-type (WT) Lr, but pPKC and IκBα were not decreased in treatment with *ΔdgkA* Lr. Mucosal IL-6 and systemic interleukin (IL)-1α, eotaxin, and granulocyte colony-stimulating factor (G-CSF) were suppressed in WT Lr, but not in *ΔdgkA* Lr colonized mice. Collectively, the commensal microbe Lr may act as a “microbial antihistamine” by suppressing intestinal H1R-mediated proinflammatory responses via diminished pPKC-mediated mammalian cell signaling.

**INTRODUCTION**

Diacylglycerol kinases (DGKs) are key enzymes in mammalian lipid metabolism that phosphorylate diacylglycerol (DAG) formed by the turnover of membrane phospholipids to phosphatidic acid (PA).1,2 Many mammalian hormones, growth factors, and other cell stimuli evoke a transient increase in the amounts of cellular DAG through hydrolysis of phosphoinositides by phospholipase C3 upon G protein-coupled receptor (GPCR) activation.4 As an example, when histamine binds to a G protein-coupled histamine receptor,5 DAG is an established second messenger and an allosteric activator of protein kinase C (PKC)4,6,7 for important biological processes.3,8 In mammals, DAG is a precursor in phospholipid metabolism, but it also serves as an intracellular lipid signal that activates PKC and is involved in cell cycle regulation, cell survival, tumorigenesis, and apoptosis.1,9 DGK consumes DAG to produce PA and is a potential terminator of DAG signaling. By attenuating DAG levels, DGK may downregulate membrane localization of PKC and may terminate transient receptor-induced PKC activation, thereby inhibiting signaling pathways downstream of PKC.3

In bacteria, the head group of phosphatidylglycerol is extensively used in the biosynthesis of membrane components, and removal of the head group generates DAG.8 DAG is converted to PA by Dgk2 and recycles DAG into the cytidine diphosphate-diacylglycerol pathway for phospholipid
synthesis, thereby preventing the lethal accumulation of DAG in bacterial membranes.8 Lactobacillus reuteri 6475 (Lr) genome has been sequenced,10 and this bacterial genome contains a gene encoding diacylglycerol kinase (Dgk). Gram-positive bacteria have been documented to express soluble isofoms of Dgk.3,11 As bacterial Dgk shares 19% sequence identity with mammalian DGK,8 we speculated that Lr-derived Dgk enzyme might inhibit DAG downstream signaling in bacterial and mammalian cells.3,8

A key biogenic amine, histamine, signals via type 1 histamine receptor (H1R) to affect PKC signaling in mammalian cells. Histamine serves a primary role as a mediator of allergic inflammation.12,13 Histamine is also a neurotransmitter that plays a key role in areas of the central nervous system enriched with histaminergic neurons such as the hippocampus.5,14,15 Apart from mammals, microbes can synthesize histamine to maintain intracellular pH.16,17 One such bacterium, Lactobacillus reuteri 6475, belongs to the phylum Firmicutes and is indigenous to the gastrointestinal tracts of avian and mammalian species.18 This organism is considered to be a GRAS (generally recognized as safe) and beneficial microbe, and has been used globally as a probiotic for more than two decades. A recent pangenomic study showed that Lr 6475 strain is distinctive among gut microbes by containing a complete chromosomal hdc gene cluster (genes hdcA, hdcB, and hdcP) and the genetic capacity to convert histidine to histamine.19 Lr-derived histamine suppressed proinflammatory cytokines in intestinal epithelial cells, monocytes,17,20 and intestinal inflammation in different rodent models.21,22 However, the molecular mechanisms behind the interactions of Lr-derived histamine and the mammalian intestinal epithelium are not clearly understood.

Histamine interacts with the host using four different mammalian receptors (H1R, H2R, H3R, and H4R).23 The majority of histamine receptors in the gut are known to be H1R and H2R.24,25 H1R is known to initiate proinflammatory pathways by increasing phosphorylation of protein kinase C (pPKC).5,26–28 H2R, in contrast, suppresses inflammation by inducing protein kinase A phosphorylation5,29 or by suppressing Toll-like receptor-mediated inflammation.30 H2R antagonists increased infection risk and necrotizing enterocolitis in low birth weight infants31,32 and doubled the risk of hospitalization and surgery for patients with Crohn’s disease.33 Suppression of intestinal inflammation depended on the presence of dietary l-histidine and histamine-generating Lr 6475 in a murine colitis model via a H2R-mediated mechanism.34 Histidine decarboxylase (HDC) is the unique enzyme responsible for generation of a biogenic amine, histamine, via conversion of l-histidine in mammals, including humans.14,35 HDC deficiency in mice was shown to promote inflammation-associated colorectal cancer by accumulation of CD11b+Gr-1+ immature myeloid cells (IMCs) in bone marrow and spleen compared with wild-type (WT) mice.14 This deficiency was associated with increased concentrations of proinflammatory cytokines, especially interleukin (IL)-6 and IL-1, and excessive quantities of IL-1 were generated at least partly by immature myeloid cells in the circulation.14 Histamine plays a key role in the maturation of IMCs via HDC expression.36

Histamine derived from Lr differentially promotes H2R pathway signaling and may antagonize H1R signaling in parallel. Microbiome-generated Dgk synthesized by Lr may interact with host epithelial DAG and inhibit H1R downstream signaling by converting DAG to PA, thereby reducing quantities of DAG, preventing PKC phosphorylation, and likely reducing intestinal inflammation. To better understand the molecular interactions of a single commensal bacterium Lr with the intestinal epithelium and its impact on shaping gut immune maturation, the mammalian intestine was colonized with WT or mutant (AhdcA or AdgkA) Lr strains deficient in diacylglycerol kinase or histamine production using inbred and outbred germ-free (GF) mouse models. The results described in this study show that bacterial Dgk is secreted into the extracellular milieu and modulates PKC signaling in mouse and human intestinal epithelial cells. Furthermore, microbial DAG kinase appears to modulate myeloid cell maturation and cytokine production in vivo.

RESULTS

WT and mutant L. reuteri 6475 stably colonize the GF mouse intestine

The 10-week-old BALB/c GF mice were either colonized with L. reuteri 6475 WT (WT) or AhdcA L. reuteri 6475 (AhdcA) or AdgkA L. reuteri 6475 (AdgkA) on day 1 and maintained for 22 days (Figure 1a). On day 23, the mice were killed and intestinal contents were collected and processed as described. Extracted DNA was quantified using quantitative PCR. GF mice colonized for 22 days with WT or AhdcA or AdgkA Lr were successfully colonized with a single Lr administration. We detected log108.56 ± 0.72 of WT Lr, log107.64 ± 0.76 of AhdcA, and log107.06 ± 0.66 of AdgkA per gram of cecal contents of BALB/c gnotobiotic mice (Figure 1b). Similarly, 13-week-old Swiss Webster (SW) GF mice were stably colonized with L. reuteri 6475 WT (WT) for 22 days (Supplementary Figure S1a online). However, SW GF mice yielded more abundant gut lactobacilli following colonization by Lr compared with BALB/c GF mice. On day 23, the SW mice were killed, and were colonized with log109.83 ± 0.24 of WT Lr per gram of cecal contents (Supplementary Figure S1b).

Systemic immune responses were suppressed by intestinal colonization with histamine-generating, DAG kinase-producing WT L. reuteri

To identify whether hdcA+dgkA+ Lr colonization had an impact on immunomodulation of the mouse intestine, proinflammatory cytokines including IL-1β, IL-6, tumor necrosis factor-α, IL-17, IL-22, IL-12, etoxin, granulocyte colony-stimulating factor (G-CSF), and interferon-γ were analyzed in the blood plasma of gnotobiotic and GF control mice. Interestingly, mice colonized with WT L. reuteri yielded reduced circulating concentrations of IL-1β compared with mice colonized by L. reuteri AdgkA or control GF mice.
(Figure 2a). Similar differences were observed for the chemokines, eotaxin and G-CSF, in peripheral blood quantified in BALB/c gnotobiotic mice colonized by WT **L. reuteri** and respective mutants (Figure 2b,c). In addition, circulating quantities of IL-6 were reduced in SW (outbred) mice that received WT **L. reuteri** (Supplementary Figure S2e) by orogastric gavage. In addition, amounts of plasma IL-10, a cytokine known to suppress inflammation, were increased in mice colonized by WT **L. reuteri** (Supplementary Figure S2a–f), whereas G-CSF and IL-6 were diminished (Supplementary Figure S2a,c) in SW mice colonized with WT **L. reuteri** (Supplementary Figure S2a–f).

**Intestinal mucosal proinflammatory cytokine suppression by hdcA<sup>+</sup> dgkA<sup>+</sup> WT L. reuteri in gnotobiotic mice**

Mucosal immune responses because of the presence and absence of histamine and/or diacylglycerol kinase of WT **L. reuteri** 6475 were measured in intestinal mucosal tissue samples. Cecal IL-6 mRNA in the mucosa was significantly decreased in mice colonized by WT **L. reuteri** as compared with mice colonized by mutant AdgkA **L. reuteri** or control GF BALB/c mice (Figure 2d). However, other cytokine mRNAs such as tumor necrosis factor-α, interferon-γ, IL-17, IL-12, IL-1x, and IL-22 were not altered or below the detection limit. Similarly, IL-6 mRNA and protein quantities were significantly

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**Figure 1** Colonization status of histamine-generating probiotic strain (**Lactobacillus reuteri**/6475) in adult BALB/c mice. (a) BALB/c mice (10 weeks old) were colonized with **L. reuteri** (Lr) 6475 on day 1 and killed on day 23. The mice were fed with a histidine-rich (4%) diet during the bacterial colonization period. Four groups differed by probiotic colonization status: group 1 germ-free (GF) control mice that lacked bacteria, group 2 mice received wild-type (WT) **L. reuteri** 6475, group 3 received ΔhdcA **L. reuteri** 6475, and group 4 received ΔdgkA **L. reuteri** 6475 (n = 10 mice per group, 5 males and 5 females). (b) Cecal intestinal contents showed positive bacterial colonization of WT, ΔhdcA, and ΔdgkA Lr after day 22 days of colonization. GF mice showed no amplification indicating the germ-free status. Data are expressed as mean ± s.e.m. *P* < 0.05 and ***P* < 0.001; n = 10 mice per group. One-way analysis of variance with Bonferroni correction. (c) Transmission electron microscopy or (c) Scanning electron microscopy of HT29 cells co-cultured with WT Lr 6475 for (l) 2 h or (ll) 6 h under anoxic condition. A full color version of this figure is available at the Mucosal Immunology journal online.
reduced in the cecal mucosa of SW mice colonized by WT L. reuteri compared with control GF mice (Supplementary Figure S2g,i,j).

Colonization by WT *dgkA*+, but not mutant *AdgkA*, *L. reuteri* inhibits mucosal PKC phosphorylation and NF-κB accumulation in intestinal epithelium

As PKC phosphorylation depends on availability of DAG through GPCR activation,5 phosphorylation of pPKC was compared among the cecal mucosae of mice colonized by WT and mutant *ΔhdcA* or *AdgkA L. reuteri*, in addition to control GF mice. To determine that the bacterial enzyme Dgk, presumably released into the extracellular milieu, inhibits DAG-mediated signaling and PKC phosphorylation in mammalian cells, immunoblot studies were performed targeting PKC phosphorylation using specific antibodies. The phosphorylated protein kinase pPKCα (Ser 657) was diminished in the cecal mucosa of gnotobiotic BALB/c mice colonized with WT Lr compared with control GF mice. In addition, BALB/c mice colonized by mutant *ΔhdcA L. reuteri* had reduced amounts of pPKC when compared with control GF mice (Figure 3a). Interestingly, the greatest abundance of pPKC was localized in the intestines of mice colonized by mutant *AdgkA L. reuteri* as compared with GF mice colonized by WT L. reuteri or mutant *ΔhdcA L. reuteri* (Figure 3a,c). Consistent with the immunoblot data, pPKC was decreased in ceca of mice colonized by WT and mutant *ΔhdcA L. reuteri* when compared with mice colonized by mutant *AdgkA L. reuteri* (Figure 3b) evaluated by fluorescence immunohistochemistry. SW GF mice colonized by WT L. reuteri also showed reduced levels of pPKC compared with corresponding nonphosphorylated PKC and GF PKC protein concentrations from the cecal mucosa (Supplementary Figure S3a).

Human jejunal enteroids treated with WT *L. reuteri* conditioned media (CM) showed decreased pPKC levels compared with human enteroids treated with mutant *AdgkA L. reuteri* CM (Supplementary Figure S4a). In addition, *ΔhdcA Lr* CM-treated enteroids showed a trend toward increased PKC phosphorylation (Supplementary Figure S4a). Consistent with immunoblot data, pPKC was decreased in human enteroids treated with WT *L. reuteri* CM compared with mutant *AdgkA Lr* CM, as examined by fluorescence immunohistochemistry (Supplementary Figure S4b). To explore mammalian intestinal epithelial DAG and pPKC signaling...
further, ileal enteroids from 10-week-old GF mice were treated with a DGK inhibitor in vitro. The enteroids treated with a DGK inhibitor (R59-022) yielded increased PKC phosphorylation in the presence of WT Lr CM, whereas enteroids lacking DGK inhibitor in the presence of WT Lr CM did not yield evidence of increase PKC phosphorylation (Supplementary Figure S5). Similarly, in vitro treatment of HT-29 human epithelial cells with \( \Delta dgkA \) Lr CM yielded a nearly threefold increase in PKC phosphorylation compared with treatment with Lr WT CM (Supplementary Figure S4c). In addition, nonphosphorylated

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**Figure 3**  *Lactobacillus reuteri* derived diacylglycerol kinase (Dgk) synthesis prevents type 1 histamine receptor (H1R)-mediated protein kinase C (PKC) phosphorylation and nuclear factor (NF)-κB accumulation in mammalian intestinal epithelium. (a) BALB/c mice (10 weeks old) colonized with *L. reuteri* (Lr) 6475 on day 1 and killed on day 23 showed suppressed cecal pPKC. Western blot analysis of cecal mucosa proteins from wild-type (WT), \( \Delta hdcA \), and \( \Delta dgkA \) Lr colonized mice and germ-free (GF) mice incubated with antibody targeting phospho-PKC (pPKC) or non-phospho-PKC (PKC). (1) Proteins pooled from all the five female mice and (2) protein pooled from all the five male mice per group. The proteins were normalized for housekeeping protein α-tubulin. The pPKC and PKC ratio was obtained by ImageJ analysis where pPKC and PKC from GF control mice was set to 1 and used as baseline. Then, 25 μg of proteins were loaded into each wells of the sodium dodecyl sulfate (SDS) gel. Proteins were isolated from the pooled samples (1 to 5 = 1 and 6 to 10 = 2) 3 individual times and proteins were estimated by Bio-Rad analysis and western blot was performed for 3 individual times. (b) The pPKC and PKC analyzed using fluorescence immunohistochemistry from cecum tissue of BALB/c mice colonized with WT, \( \Delta hdcA \), and \( \Delta dgkA \) L. *reuteri*6475, and control GF mice. Red color indicates pPKC, pink color indicates PKC distribution of lamina propria, and green color indicates nuclei staining by 4',6-diamidino-2-phenylindole (DAPI). Bar = 10 μm. Images shown are representative of five individual mouse sections per group. The proteins were normalized for housekeeping protein α-tubulin. (a, b) Blue arrow indicates downregulation of pPKC and yellow arrow indicates upregulation of pPKC. (c) Mean values of three independent western blot gels showing pPKC protein concentration (loaded 25 μg) and quantified by densitometry using ImageJ. GF control groups were set to 1. ***P < 0.001; n = 10 mice per group. One-way analysis of variance with Bonferroni correction. (d) Western blot analysis of cecal mucosa proteins from WT, \( \Delta hdcA \), and \( \Delta dgkA \) Lr colonized, and GF mice incubated with antibody targeting IκBα (L35A5) NF-κB complex.
PKC protein concentrations increased over time in the presence of WT Lr CM and decreased with mutant AdgkA Lr CM treatment in HT-29 human epithelial cells (Supplementary Figure S4c).

Activation and accumulation of cytosolic nuclear factor (NF)-κB (IkBz-RelA:P50 complex) is dependent on phosphorylation of pPKC. The intestinal epithelial tissues obtained from the cecal mucosa of mice colonized by WT, mutant AhdcA, or AdgkA L. reuteri were compared with control GF mice. To determine that the bacterial enzyme Dgk inhibits activation of NF-κB through reduced phosphorylation of PKC in mammalian cells, immunoblot studies were performed targeting NF-κB accumulation using specific antibodies. Total IkBz (L35A5) complex was diminished in the cecal mucosa of gnotobiotic BALB/c mice colonized with WT Lr as compared with control GF mice. The AdgkA L. reuteri colonized mice yielded increased synthesis of NF-κB (Figure 3d).

**L. reuteri** DAG kinase is secreted into the extracellular milieu

As we observed negligible PKC phosphorylation in intestinal mucosal tissues of mice colonized with WT or AhdcA compared with AdgkA Lr, Dgk (~15 kDa) is secreted by the bacterium Lr, and the extracellular microbial enzyme may effectively decrease DAG activity in mammalian cells. By suppressing DAG lipid-mediated signaling, Dgk kinase from lactobacilli may suppress H1R signaling via reduced PKC phosphorylation. To explore the possibility that bacterial Dgk may be secreted into the intestinal lumen and interact with substrate DAG in the host epithelium, Dgk was detected in bacterial culture supernatants following growth in liquid media. WT and mutant hdcA or AdgkA L. reuteri 6475 were cultured in LDM4 for 12 h in the presence and absence of DAG, and liquid culture supernatants were evaluated.

Extracellular proteins were fractionated by size, processed for tryptic digestion, and analyzed by liquid chromatography/tandem-mass spectrometry followed by amino acid sequence analysis. Amino acid sequences of Dgk peptide fragments (Supplementary Figure S6) matched the expected Dgk protein sequences (Figure 4a–c). As expected, Dgk was present in WT L. reuteri supernatants, but absent in mutant AdgkA L. reuteri cultures (Figure 4c). WT Lr dgkA mRNA levels were significantly increased in the presence of DAG compared with absence of DAG in the growth medium after 12 h in vitro (Figure 4d) and bacterial dgkA gene expression was increased in cecal luminal contents of the gnotobiotic BALB/c mouse intestine colonized by WT L. reuteri (Figure 4f). In addition, clade II L. reuteri 6475 dgkA gene was present in greatest quantities during the elongation phase (Supplementary Figure S7). Bacterial mRNA was obtained from the cecal contents and quantified using reverse transcriptase-PCR. Bacterial dgkA expression was increased in the lumens of both WT and AhdcA Lr colonized gnotobiotic BALB/c mice compared with AdgkA Lr colonized mice, whereas GF control BALB/c mice showed no evidence of dgkA expression (Figure 4f). In addition, mammalian DGK quantified from the cecal intestinal epithelium by immunoblot did not yield significant differences between control mice and the GF mice colonized with WT or mutant L. reuteri (Supplementary Figure S3b).

**Bacterial histidine decarboxylase mRNA and luminal histamine present in the intestinal lumen of gnotobiotic BALB/c mice colonized by L. reuteri**

Luminal contents were treated with 99% methanol and the supernatants obtained were used for histamine quantification using liquid chromatography/mass spectrometry. WT L. reuteri 6475 and mutant AdgkA L. reuteri colonized BALB/c mice yielded detectable luminal histamine (qualitative data not shown). Interestingly AdgkA Lr culture grown for 24 h in lactobacillus defined media, LDM3 or LDM4, generated more than twofold increased histamine concentrations compared with WT Lr culture. Mutant AhdcA Lr yielded absence of detectable histamine in liquid media (Supplementary Table S1). However, mutant AhdcA Lr lacked detectable histamine in culture supernatants (Supplementary Table S1). Bacterial histidine decarboxylase mRNA, presumably at least partly from L. reuteri, was also detectable in the intestinal lumens of gnotobiotic BALB/c mice colonized by WT and mutant AdgkA L. reuteri (Figure 4e).

**L. reuteri** resides adjacent to the intestinal epithelium and modifies the intestinal mucus layer and GPCR signal of gnotobiotic mice

Bacterial fluorescent in situ hybridization demonstrated that Lr resides in the mucus layer, penetrates the inner mucus layer, and establishes intimate contact with intestinal enterocytes in the murine large intestine (Figure 5a–f). The data suggest that Lr directly communicates with the host intestinal epithelium irrespective of WT or mutant (hdcA, dgkA) status. Similarly, Lr was adjacent to the human epithelium when Lr was co-cultured with human jejunal enteroid monolayers (Figure 5g,h) and HT-29 cells (Figure 1c).

To determine H1R and H2R GPCR protein quantities, fluorescence immunohistochemistry analysis was performed as described in the Methods. Both H1R and H2R protein expression were similar between the WT, AhdcA, and AdgkA Lr colonized mice compared with control GF mice (Figure 5i–p). However, H1R was present in greater abundance than H2R in the intestinal epithelium.

Mucus serves as the primary contact between the intestinal epithelial layer and luminal antigens. As GF mice were colonized with Lr, it was of interest to evaluate mucus maturation after probiotic colonization using lectin staining. We observed significantly increased mucus fucosylation with Ulex europaeus agglutinin staining of the terminal mucus fucose in WT Lr colonized compared with control BALB/c GF mice (Supplementary Figure S8ai and aii). Similarly, increased fucosylation was observed in mice colonized with AhdcA and AdgkA Lr compared with control BALB/c GF mice. However, AdgkA colonized mice had decreased mucus fucosylation compared with WT Lr colonized mice (Supplementary Figure S8ai–aiv). Similarly, SW mice colonized with WT Lr also showed increased mucus fucosylation with increased
Figure 4  *Lactobacillus reuteri* 6475 derived diacylglycerol kinase (Dgk) secreted into the medium. (a) Conditioned media (CM) from wild-type (WT) *L. reuteri* (Lr) 6475 grown in LDM4 with lipid diacylglycerol (DAG) supplements for 12 h were analyzed using Eksigent nanoLC and the ABCIEX TripleTOF 5600 mass spectrometer. The obtained sequences were matched with the NCBI database with the Analyst software version 1.6 (ABCIEX). Red line indicates the protein sequence matching *L. reuteri* Dgk protein in the database from medium grown in the presence of lipid DAG, whereas blue line indicates the protein sequence matching *L. reuteri* Dgk in the database from medium grown in the absence of lipid DAG. The protein sequences were obtained from 20 kDa band (also (c)). (b) CM from WT, *dhcA*, and *dgkA* Lr 6475 grown in LDM4 without lipid DAG supplement for 12 h were analyzed as mentioned in a. (c) One-dimensional sodium dodecyl sulfate (SDS) gel showing positive bands at 20 and 10 kDa from *L. reuteri* derived CM with and without DAG supplementation. (a–c) The 20 kDa protein band was cut and followed by tryptic digestion and were used to analyze as mentioned in a and b. (d) mRNA expression of *dgkA* in WT *L. reuteri* cultured for 12 h with and without lipid DAG supplementation in LDM4 minimal media in N2/CO2 at 37 °C and analyzed by reverse transcriptase-PCR (RT-PCR). All experiments were repeated 3 times individually; *n* = 3 per group. Data are expressed as mean ± s.e.m. *P* < 0.05, one-way analysis of variance with Bonferroni correction. Graph showing cecal bacterial (e) *dhcA* and (f) *dgkA* gene expression levels of BALB/c mice colonized with WT, *dhcA*, *dgkA* Lr, and germ-free (GF) control. (e, f) Data are expressed as median with range. *P* < 0.05; *n* = 10 mice per group. Kruskal–Wallis test with Dunn’s correction. A full color version of this figure is available at the *Mucosal Immunology* journal online.
mucus secreting goblet cells (Supplementary Figure S8av–aviii and 8b).

Histamine-dependent enhancement of CD11b+ Gr-1+ Ly6Ghigh splenic myeloid cell maturation was offset by the absence of DAG kinase in L. reuteri. To determine whether bacterial-derived histamine has immunomodulatory effects by targeting myeloid cell maturation, GF mice were colonized with WT, ΔhdcA, ΔdgkA Lr, and a control group without any bacteria. The absence of endogenous histamine results in increased proportions of IMCs and has been associated with cancer progression in mammals.14 Azoxymethane/dextran sodium sulfate-treated Hdc−/− male mice showed significantly reduced CD11b+ Gr-1+ IMCs (80%) in spleens after oral administration of hdc+ Lr 6475 (WT) treatment.41 In this study, BALB/c gnotobiotic mice

Figure 5. Lactobacillus reuteri resides close to intestinal epithelium and epithelial histamine receptors. Fluorescent in situ hybridization (FISH) analysis used to see L. reuteri (Lr) localization in the mouse intestine (a–c) wild-type (WT), (d, e) ΔhdcA, and (f) ΔdgkA Lr 6475. Red color indicates the bacterium and blue color (or) white color indicates the nuclei stained with 4',6-diamidino-2-phenylindole (DAPI). Original magnification ×100. Representative images of six mice per group. Bar = 200 pixels. (g, h) WT L. reuteri 6475 interaction on the human jejunal enteroids monolayer visualized by FISH. Red color indicates the bacterium and blue color (or) white color indicates the nuclei stained with DAPI. Original magnification ×100. Bar = 200 pixels. L, luminal; S, serosal side. Histamine receptor 1 (H1R) analyzed using fluorescence immunohistochemistry from cecum tissue of 10-week-old (i) germ-free (GF) BALB/c mice colonized with (j) WT, (k) ΔhdcA, (l) and ΔdgkA Lr 6475. Green color indicates H1R distribution of cecal epithelium and blue color indicates nuclei by DAPI staining. Histamine receptor 2 (H2R) analyzed using fluorescence immunohistochemistry from cecum tissue of 13-week-old (m) GF BALB/c mice colonized with (n) WT, (o) ΔhdcA, and (p) ΔdgkA L. reuteri 6475. Red color indicates H2R distribution and blue color shows nuclei by DAPI staining. (i–p) Original magnification ×60. Bar = 10 μm. Representative images of six mice per group.
Lactobacillus reuteri colonization reduces splenic immature myeloid cells (IMCs) with increased cecal mmu-miR-669k-3p. BALB/c mice (10 weeks old) were colonized with wild-type (WT), ΔhdCA, ΔdgkA L. reuteri (Lr) 6475, and germ-free (GF) control. (a) Zebra plot showing CD11b⁺ Gr1⁺ (a*) IMCs with (a*) Ly6G⁹⁺, Ly6G⁸⁺, and Ly6G⁷⁺ in spleen of mice. Representative images of n = 10 mice per group; hi, high; lo, low; med, medium. (b) Splenic IMCs of WT, ΔhdCA, and ΔdgkA Lr colonized, and control GF mice. Data are expressed as mean ± s.e.m. *P < 0.05; n = 10 mice per group. One-way analysis of variance with Bonferroni correction. (c) Table shows microRNA (miRNA) data obtained from cecum mucosa of WT, ΔhdCA, and ΔdgkA Lr colonized, and control GF mice quantified using ViiA7 real-time reverse transcriptase-PCR (rtRT-PCR). From 84 miRNAs quantified, one miRNA, mmu-miR-669k-3p (**), showed significant increase. Each group represents cecum mucosal miRNA pooled from 10 mice. Significant differences and ΔCt were obtained by web-based BmiRNA PCR Array Data Analysis software (QIAGEN). A full color version of this figure is available at the Mucosal Immunology journal online.
colonized with WT *L. reuteri* strain yielded significantly decreased proportions of CD11b⁺Gr-1⁺ IMCs in the spleen (Figure 6b), suggesting that histamine-generating *Lr* promotes maturation of splenic CD11b⁺Gr-1⁺ IMCs (Figure 6a,b). The lack of bacterial histidine decarboxylase in ΔhdcA *Lr* corresponded with a trend toward increased populations of IMCs in the spleen (Figure 6b). In addition, GF mice colonized with ΔdgkA *Lr* did not result in suppression of IMCs in the spleen (Figure 6b), suggesting that loss of microbiome-derived DAG kinase and enhanced H1R signaling may offset effects via H2R on myeloid cell maturation. In addition, we observed reduced Ly6Ghigh and Ly6Gmedium populations in the spleens of mice that were colonized with WT *L. reuteri* as compared with mice colonized by mutant ΔhdcA or ΔdgkA *L. reuteri* (Figure 6a).

**L. reuteri** colonization leads to increased microRNA mmu-miR-669k-3p in the intestinal epithelium of GF BALB/c mice

Because basal cytokine responses were suppressed in the presence of WT *L. reuteri* in the intestines of gnotobiotic BALB/c mice (Figure 2), changes in inflammatory responses were related to microRNA modification. A total of 84 individual microRNAs considered as candidates for regulating inflammatory pathways were selected. The microRNA expression patterns in the intestinal mucosa were evaluated for alterations by *Lr*-derived histamine and/or Dgk. The miRNA mmu-miR-669k-3p was found to be increased by twofold in mice colonized with WT *Lr* compared with ΔhdcA, ΔdgkA *Lr*, and control GF mice (Figure 6d).

**DISCUSSION**

The mammalian microbiome contains microbial cells with genomes encoding the capacity for microbial-mediated conversion of amino acids to bioactive signals such as biogenic amines. As shown in this study, bacterial enzymes such as diacylglycerol kinase may abrogate mammalian cell signaling pathways by effectively “removing” mammalian lipid signals such as diacylglycerol. The gut commensal microbe, *L. reuteri*, sequesters Dgk and because of its close proximity to the gut epithelium, this enzyme may convert mammalian DAG to PA. The presence of dgkA-producing *L. reuteri* suppresses phosphorylation of PKC downstream of the histamine type 1 receptor. Dgk secreted by *Lr* 6475 potentially interacts with epithelial DAG lipid signaling, thereby inhibiting proinflammatory H1R and allowing “net positive” activation of H2R by histamine-generating *Lr*. Ultimately, this reduced phosphorylation of PKC may result in diminished signaling via NF-κB and suppression of mucosal inflammation.

Human-derived clade II *L. reuteri* strains contain a complete chromosomal hdc gene cluster with genes encoding histidine decarboxylase (HdcA) and the histidine/histamine transporter (HdcP), culminating in suppression of human tumor necrosis factor production. Gut microbes like *L. reuteri* clade II strains provide a useful model system to study the effects of luminal histamine on the mammalian gut mucosa. By introducing individual gut microbes into GF mice, our data show that WT and mutant *L. reuteri* each colonized the mammalian intestine successfully with a single oral dosing. Although immunologists have accumulated insights regarding how histamine works in allergic inflammation and in the context of mucosal mast cells, the role of histamine in the lumen of the gastrointestinal tract is not well understood. The effects of luminal histamine may differ from that of mucosal histamine simply based on the relative distributions of histamine receptors in different cell types. Our studies suggest that luminal histamine does affect mucosal and systemic immunity, as well as mammalian epithelial cell signaling pathways. Suppression of IL-6 production within the cecal mucosa is one example of histamine-dependent suppression of gut inflammation in an acute colitis model. By investigating the role of luminal histamine produced by gut microbes, we may gain a deeper mechanistic understanding of the interrelationships between the gut microbiome, the intestinal mucosa, and immunity.

Luminal histamine may act through the histamine type 2 receptors (to suppress the production of proinflammatory cytokines and cell signaling via mitogen-activated protein kinases (see Thomas et al. and Gao et al.)). However, both H1R and H2R are present in the mouse and human intestinal epithelium, and hence the role of H1R in promoting gastrointestinal mucosal inflammation must be considered. Our studies provide insights into a potential “brake” directed at H1R-mediated signaling. A bacterial enzyme, Dgk, may modify the lipid signal diacylglycerol to abrogate H1 receptor-mediated signaling via PKC. In contrast, histamine binding to H2R activates protein kinase A and binds cAMP via adenylyl cyclase to suppress mitogen-activated protein kinase activation. Instead, H1R signaling involves the conversion of phosphatidylinositol 2 by phospholipase C into DAG and inositol trisphosphate. DAG stimulates phosphorylation of PKC by binding to the catalytic domain 1 and recruiting PKC to the membrane. The interaction of PKC with the membrane causes release of pseudosubstrate from the catalytic domain leading to availability of serine and threonine phosphoacceptor residues causing PKC activation. Phosphorylated PKCs (pPKC) is known to induce proinflammatory responses such as NF-κB activation. Thus, a net reduction of pPKC in the intestinal mucosa by the gut microbiome may help to keep the immune system “in check” and promote immunologic homeostasis.

Beneficial microbes in the intestine may produce enzymes that inhibit mammalian cell signaling pathways by effective signal depletion. DGK is an enzyme known to inhibit DAG-mediated signaling by converting DAG to (PA utilizing adenosine triphosphate as the phosphate source. Conversion of DAG to PA depletes DAG, and this reduced amount of DAG may suppress PKC activity, thereby suppressing proinflammatory immune responses in the intestine (Figure 7). On that note, *Escherichia coli* has increased amounts of DAG kinase. Resting human platelets (10⁹ cells have 0.111 nmol DAG) treated with *E. coli* DAG kinase results in conversion of lipid DAG to PA. We propose that the mammalian microbiome may synthesize Dgk adjacent to the
intestinal epithelium, effectively depleting mammalian DAG and inhibiting H1R downstream activation. In support of our proposed hypothesis, *L. reuteri* is positioned adjacent to the intestinal epithelium and could deliver DAG kinase in close proximity to the epithelial cell membranes. PKC phosphorylation was drastically reduced in the intestinal epithelium of WT compared with *ΔdgkA* Lr colonized mice. Dramatically increased phosphorylation of PKC was evident in the intestinal mucosa of mice colonized by *ΔdgkA* Lr and control GF BALB/c mice, but not in mice colonized by WT or *AhdcA L. reuteri*. Increased PKC phosphorylation was observed in human cells such as the human intestinal epithelial cell line HT29 treated with *ΔdgkA* Lr-CM. These findings raise intriguing possibilities of bacterial enzymes from the mammalian microbiome converting compounds within the mammalian cell milieu and altering signaling pathways in mammalian cells. Mammalian epithelial cells may, for brief periods of time, behave like "chimeric" cells with bacterial and mammalian enzymes affecting intra-mammalian cell signaling pathways.

Previous findings showed that bacterial Dgk shares 18 to 19% amino acid sequence identity with human DGK\(^*\) that can have soluble isoforms.\(^{11,38}\) Similarly, *Staphylococcus aureus* DAG kinase B structure has been identified to be directly relevant to the human enzymes that function in cell signaling because the key structural and active site residues are conserved in the sequences of both enzymes.\(^5\) The bacterial Dgk superfamily provides candidate enzymes and substrates for evaluation of small-molecule therapeutics to combat Gram-positive organisms and to modulate lipid signaling pathways in humans.\(^8\) Mammalian DGK is found both in soluble and membrane-bound forms actively converting lipid DAG to PA,\(^{31}\) thereby suppressing PKC phosphorylation via DAG phosphorylation to PA.\(^{32}\) We hypothesized that Dgk released by Dgk-positive intestinal lactobacilli can suppress H1R downstream signaling by modifying mammalian cell membrane-associated DAG. It is not understood whether a bacterial kinase from the microbiome may directly convert membrane-associated DAG in adjacent mammalian epithelial cells. We present evidence that downstream mammalian PKC is activated in the absence of gut microbes, but suppressed in the presence of histamine-generating microbes with an intact DAG kinase gene. In support of the hypothesis, Lr-derived Dgk could not suppress intestinal epithelial PKC phosphorylation in the presence of a DGK inhibitor.

Prior evidence indicates that bacterial DAG kinase may be secreted or released by microbes into the extracellular milieu.\(^8,50\) Prior studies have also indicated that mammalian DAG kinases may be released;\(^{51}\) however, mammalian DAG kinase was not significantly altered by the presence of *L. reuteri* DAG kinase in gnotobiotic mice. DAG kinases bind to lipid DAG and generate PA in the lipid bilayer and these compounds may be available for effects on intracellular signaling pathways. *Bacillus subtilis* has been shown to produce 34 kDa soluble Dgk enzyme.\(^11\) We looked for soluble Dgk proteins/peptides in the supernatant obtained from WT Lr culture for 12 h in LDM4. By liquid chromatography/tandem-mass spectrometry analysis we generated the presence of six peptide sequences from bacterial culture supernatants, and these peptide sequences were identical to segments of the Dgk protein (http://www.ncbi.nlm.nih.gov/protein/EGC15283.1) from Lr 6475. In addition, Lr 6475 was found to colonize the germ-free mouse intestine, adjacent to the intestinal epithelium. *L. reuteri* is known to possess secretory and extracellular proteins (secretome) and these secreted proteins include enzymes involved in lipid metabolism.\(^{53,54}\) DAG is a well-known mammalian membrane lipid compound that flip-flops rapidly in the lipid bilayer.\(^{55}\) Dgk secreted by *L. reuteri* may interact with host DAG in the gut epithelium, reducing amounts of DAG and thereby restricting PKC phosphorylation and NF-kB activation following downregulation of mucosal cytokines like IL-6. Such a reduction in mucosal inflammation or dampening of mucosal immunity may result in altered maturation programs of splenic myeloid cells.\(^{14}\)

A deficiency HDC and lack of endogenous histamine was shown to promote inflammation-associated colorectal cancer by accumulation of CD11b \(^+\) Gr-1 \(^+\) IMCs.\(^{14,56}\) Modulation of histamine receptors through its antagonist modifies colorectal cancer progression.\(^{57–59}\) We observed suppression of splenic (not bone marrow) IMCs in germ-free mice colonized with WT *L. reuteri*. This suppression could be due to reduced IL-6 expression in intestinal mucosa because increased IL-6 was previously shown to increase splenic IMC infiltration.\(^{14,60}\) Suppression of IL-6 facilitates maturation of IMCs.\(^{14,36}\)
However, AdgkA Lr colonized mice did not suppress splenic IMCs or reduce IL-6 mRNA expression, suggesting that dgkA is important in alteration of H1R-mediated signaling and maturation of myeloid cells in peripheral compartments. Bacterial genes hdcA and dgkA are important for L. reuteri to generate sustained immunosuppressive effects in the intestine. In addition to reduced IL-6 gene levels we also observed reduced IL-1α in blood plasma of mice that received WT L. reuteri compared with the GF control mice. Interestingly, AdgkA Lr colonized mice had increased levels of plasma IL-1α, a proinflammatory cytokine predominantly secreted by macrophages. IL-1α may promote IL-6 production. As we observed decreased IL-1α protein levels in blood plasma of WT Lr colonized mice but not in AhdcA and AdgkA Lr colonized mice, we speculate that luminal histamine and luminal DAG kinase may both be important for IL-1α production by mucosal macrophages. Furthermore, we observed diminished mucosal IL-6 mRNA levels in the presence of WT Lr 6475 with increase in microRNA (mmu-miR)-669k-3p of BALB/c mice. Therefore, we speculate that mmu-miR-669k-3p expression modulates IL-6 gene expression and may have an impact on myeloid cell maturation. However, detailed studies are required to further characterize the microRNA mmu-miR-669-k-3p on intestinal immunomodulation.

The mammalian microbiome includes dozens of bacterial taxa that possess the genetic capacity to perform intestinal luminal conversion of diverse biological compounds. These compounds may be derived from the diet or mammalian cells and provide substrates for bacterial enzymes. The model commensal microbe, L. reuteri, has yielded insights regarding luminal conversion by gut microbes and effects on epithelial and immune cell signaling pathways. Gut lactobacilli may generate small amines like histamine from amino acids and suppress inflammation via specific receptors. Gut microbes may also provide enzymes that convert lipid signals, thereby resulting in diminished signaling via receptors that promote allergic inflammation. The net effect is suppression of intestinal inflammation. Future advances in microbiome science may lead to new therapeutics that target bacterial enzymes acting in bacterial cells or mammalian cells (or both cell types) to alter cell signal or immune responses. Probiotics generating “microbial antihistamine factors” like DAG kinase may abrogate intestinal inflammation and promote the development of new microbiome-harvested treatment strategies.

METHODS

Bacterial strains. The bacterial strains used in this study were L. reuteri 6475 (or also known as MM4-1A), AhdcA Lr 6475, and AdgkA Lr 6475. All strains were routinely cultured at 37 °C in MRS (Man, Rogosa, and Sharpe) or LDM4 media and incubated overnight under strictly anoxic conditions using N2/CO2 (80:20, v/v) as the gas phase.

Gnotobiotic mouse model: bacterial colonization studies. GF BALB/c mice (five males and five females per group) were bred in Trexler-type isolators. After weaning, all mice (10 weeks old) were colonized by gavaging with L. reuteri 6475 (WT strain), AhdcA Lr 6475 (hdcA mutant strain), or AdgkA Lr 6475 (dgkA mutant strain) (1 × 10^9 bacterial cells) in the presence of a 4% L-histidine diet on day 1 and were killed on day 23. All mouse experiments were performed in a germ-free animal facility, according to an institutional animal care and use committee-approved mouse protocol at Baylor College of Medicine Houston, TX, and the Massachusetts Institute of Technology Committee on Animal Care (for elaborate protocols see Supplementary Text).

Immunoblot studies of pPKC signaling. Proteins were extracted from tissue samples in RIPA buffer. A total of 30 or 20 μg of protein from each sample was separated by sodium dodecyl sulfate–polyacrylamide gel electrophoresis and transferred onto nitrocellulose membranes. Membranes were incubated with specific primary antibodies, nonphosphorylated PKCα and pPKCα (Ser 657) (diluted 1:1,000) (Santa Cruz Biotechnology, Santa Cruz, CA). Statistics. Data were tested for normal distribution using the Kolmogorov–Smirnov test. Normally distributed data are presented as means with s.e., whereas the medians with their range are given for nonnormally distributed data. Significance of differences between Control, WT Lr, AhdcA, and AdgkA mutant Lr colonized BALB/c mice were analyzed using the one-way analysis of variance test for normally distributed data (or) the Kruskal–Wallis test for nonnormally distributed data, followed by either Bonferroni/Tukey or Dunn’s comparison post hoc tests. Differences between with and without DAG-treated bacterial culture gene expression and SW mice with and without WT Lr were analyzed using Student’s t-test followed by the Mann–Whitney test for nonnormally distributed data. Differences between the groups were considered significant at *P<0.05, **P<0.01, and ***P<0.001. SPSS 16.0 (IBM, Armonk, NY) for Windows 7 was used for data analysis. Prism 5.0 software (GraphPad Software, La Jolla, CA) for Windows was used for data presentation and also for data analysis.

SUPPLEMENTARY MATERIAL is linked to the online version of the paper at http://www.nature.com/mi

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AUTHOR CONTRIBUTIONS

B.P.G. was involved in the study design, completion of all experiments, data analysis, and interpretation and manuscript preparation. A.H. contributed to the dgkA mutant L. reuteri. S.A. helped with western blot analysis and scientific discussion. J.W.N. took part in mRNA isolation and scientific discussion. R.F. took part in flow analysis. A.H. performed histamine quantification by MS. M.E. and M.L. performed all the fluorescence immunohistochemistry staining. S.V. performed the MAGPIX assay. M.W. helped with Swiss Webster animal experimental set up at the Massachusetts Institute of Technology (MIT), Cambridge, M.W., J.G.F., and J.V. provided guidance, helped to design the experiments, and wrote the manuscript.

DISCLOSURE

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