Leuconostoc mesenteroides fermentation produces butyric acid and mediates Ffar2 to regulate blood glucose and insulin in type 1 diabetic mice

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Type 1 diabetic patients have lower counts of butyric acid-producing bacteria in the dysbiotic gut microbiome. In this study, we demonstrate that a butyric acid-producing Leuconostoc mesenteroides (L. mesenteroides) EH-1 strain isolated from Mongolian curd cheese can reduce blood glucose and IL-6 in the type 1 diabetic mouse model. L. mesenteroides EH-1 fermentation yielded high concentrations of butyric acid both in vitro and in vivo. Butyric acid or L. mesenteroides EH-1 increased the amounts of insulin in Min6 cell culture and streptozotocin (STZ)-induced diabetic mice. Inhibition or siRNA knockdown of free fatty acid receptor 2 (Ffar2) considerably reduced the anti-diabetic effect of probiotic L. mesenteroides EH-1 or butyric acid by lowering the level of blood glucose. We here demonstrate that Ffar2 mediated the effects of L. mesenteroides EH-1 and butyric acid on regulation of blood glucose and insulin in type 1 diabetic mice.

Type 1 diabetes is caused by marked insulin deficiency as a result of the loss of beta cells1–4. Hyperglycemia in type 1 diabetes probably results from a long-term imbalance between immune-mediated beta cell damage5 and beta cell repair/regeneration6. Type 1 diabetes is characterized by the presence of hyperglycemia together with insulin resistance, oxidative stress as well as elevated production of cytokines, such as C-reactive protein, interleukin (IL)-6 and tumor necrosis factor (TNF)-α7.

The role of bacteria in diabetes has been presented in animal models8. For example, the feeding of probiotic bacteria, mostly lactic acid bacteria, to diabetes-prone rats or non-obese diabetic mice can prevent or delay diabetes8–11. Probiotics can reduce blood glucose through the inflammatory attenuation and prevention of pancreatic beta cell destruction in vivo models12,13. Moreover, fermentation of bacteria in human colon and mouse cecum leads to the production of short chain fatty acids (SCFAs), such as acetate, lactate, propionate and butyrate14. The literatures found that gut microbiome-derived SCFAs also modulate different cell types in host such as pancreatic cells15, immune cells16, adipose tissue17, hepatocytes17, muscles17 and neuron cells18. Most of these cells express SCFA receptor 2 (Ffar2) and receptor 3 (Ffar3), SCFAs are detected in the blood circulation14,19–21. The evidences suggest that SCFAs can manipulate such cells to regulate the health of host.

SCFAs have been proposed as therapeutic modalities against diabetes with obesity, adipose inflammation and insulin resistance22. Notably, butyrate supplementation increases insulin sensitivity, energy expenditure23,24, and the beta cell proliferation25. Furthermore, butyrate and butyrate-producing microbes are decreased in diabetes mellitus26–28. Type 1 diabetic children have a lower relative abundance of butyrate-producing bacteria29. In our current study, we evaluate the long term effects of the oral administration of Leuconostoc mesenteroides (L. mesenteroides) EH-1 on blood glucose and insulin in type 1 diabetic mice.

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mesenteroides), a Gram-positive bacterium referred to as a L. mesenteroides EH-1 strain isolated from Aaruul or Mongolian curd cheese, supplement on the diabetic status of streptozotocin (STZ)-induced diabetic mice. We further evaluate the role of butyric acid, a fermentation metabolite of L. mesenteroides in the regulation of blood glucose in this model.

Methods

**Bacterial culture and identification.** Mongolian curd cheese was homogenized in 500 μL of sterile PBS with a grinder. Bacteria in the homogenate were plated on a tryptic soy broth (TSB) (Sigma, St. Louis, MO, USA) agar plate and incubated for 3 days at 37 °C. Sequence analysis of 16S ribosomal RNA (rRNA) genes was utilized for bacterial identification30. A single colony of bacteria from a TSB agar plate was isolated with a sterile toothpick and boiled at 100 °C for DNA extraction. Identification of L. mesenteroides EH-1 strain was validated by rRNA sequencing using the 16S rRNA 27F and 534R primers for polymerase chain reaction (PCR) (Supplementary Fig. S1)31. The 16S rRNA gene sequences were analyzed using the basic local alignment search tool (BLASTn, National Library of Medicine 8600 Rockville Pike, Bethesda, MD, USA). L. mesenteroides was cultured in TSB (Sigma) overnight at 37 °C. The cultures were diluted 1:100 and cultured to an optical density 600 nm (OD600) = 1.0. Bacteria were harvested by centrifugation at 5000 rpm for 10 min, washed with PBS, and suspended in PBS for further experiments.

**Glucose fermentation of L. mesenteroides EH-1.** To induce fermentation, L. mesenteroides EH-1 (10⁷ colony-forming unit (CFU)/mL) was incubated in rich media [10 g/L yeast extract (Biokar Diagnostics, Beauvais, France), 5 g/L TSB, 2.5 g/L K₂HPO₄ and 1.5 g/L KH₂PO₄] in the absence or presence of 20 g/L (2%) glucose at 37 °C for 24 h. Rich media or rich media plus 20 g/L glucose without bacteria were included as a control. Phenol red [0.001% (w/v), Sigma] in rich media with 20 g/L glucose served as an indicator of fermentation, converting from red-orange to yellow when fermentation occurred. High performance liquid chromatography (HPLC) was used to quantify the level of butyric acid in cultured media.

**Min6 cells treatment.** Min6 cells within 3–7 passages were cultured in Dulbecco’s modified essential medium (Gibco-BRL, Grand Island, NY, USA) with 10% (v/v) fetal bovine serum (FBS) (Irvine Scientific, Santa Ana, CA, USA). 10 mmol/L HEPES, 2 mmol/L L-glutamine, 1 mmol/L sodium pyruvate, 100 units/mL penicillin, and 100 μg/mL streptomycin. Cells were cultured for 3 days prior to analysis. After removing the media, cells were washed once with HEPES-balanced KRB (119 mmol/L NaCl, 4.74 mmol/L KCl, 2.54 mmol/L CaCl₂, 1.19 mmol/L MgCl₂, 1.19 mmol/L KH₂PO₄, 25 mmol/L NaHCO₃, 0 mmol/L HEPES, pH 7.4) containing 0.5% bovine serum albumin (BSA) without glucose. Min6 cells were preincubated for 0.5 h in 4-(2-hydroxyethyl)-1-piperazineethanesulfonic acid (HEPES)-balanced Krebs ringer buffer (KRB). After washing twice with HEPES-balanced KRB, Min6 cells were incubated for 24 h in HEPES-balanced KRB supplemented with 0.5% BSA and 100 μmol/L glucose or 100 μmol/L butyric acid or propionic acid. The media were then collected and assayed by a mouse insulin ELISA kit. For small interfering RNA (siRNA)-mediated knockdown of Ffar2, siRNA against Ffar2 was purchased from GenePharma, Shanghai, China. Min6 cells (10⁵ cells/mL) within 5–7 passages were cultured for 3 days and then reversely transfected with 10 μmol/L of Ffar2 (sense strand: 5′-GCCUGUGUGACGCUCUUAATT-3′ and anti-sense strand: 5′-UAAGAACGUCACACGCTT-3′) or scramble (sense strand: 5′-UUCCCGAAGGUCAGGCTT-3′ and anti-sense strand: 5′-ACGGUGACGGUCGGGAAT-3′) siRNAs using Lipofectamine 2000, and the media were changed 6 h thereafter. A second transfection with siRNA followed on the second day, and the experiment was performed 48 h after the second transfection as described above. RNA was extracted for quantification of Ffar2 expression by real-time PCR (RT-PCR).

**Streptozotocin (STZ)-induced type 1 diabetic mice.** The Institute Cancer Research (ICR) mice (8–12 week-old males; National Laboratory Animal Center, Taiwan) were housed at 25 °C with a 12:12 h light-dark cycle, fed a normal chow diet, and given water ad libitum. Mice (n = 4/group) were acclimatized for 5–7 days before the experiment. To induce a rapid ablation of the beta cells and hyperglycemia34 and to avoid the interruption of STZ with L. mesenteroides EH-1 or butyric acid, mice were injected with a single dose, instead of low multiple doses33, of STZ. Diabetes was induced following an 8-h fast using a single intraperitoneal (IP) injection of STZ (200 mg/kg body weight) (Sigma)32, which was dissolved in acidified citrate buffer (0.1 mol/L, pH 4.5). Two days later, after a 4-h-fast, the level of blood glucose from the tail blood was measured using a glucometer (Advantage, Roche, Mannheim, Germany). Injection of STZ for 2 days induced a weight lost (32.8 ± 0.9 vs 0.9 ± 0.9 g) and an increased blood glucose in this model.

**Diabetic mice treated with butyric acid.** Butyric acid at 4 mmol/L in water (5 mL/kg body weight) was administrated to diabetic mice via IP injection. Mice in control groups received water alone. Blood glucose was detected every day. Seven days after butyric acid injection, fasting blood was collected for detection of insulin and IL-6 as described above. For Ffar2 inhibition, a Ffar2 antagonist (GLPG-0974, Tocris Bioscience, Bristol, UK) was dissolved in dimethylsulfoxide (DMSO) to make a stock solution. Diabetic mice were injected with butyric acid (5 mL/kg body weight) taken from a stock solution of 4 mmol/L GLPG-0974 (1 mg/kg body weight) was diluted in saline then was given at 1 mL/kg body weight35 by gastric gavage just before butyric acid injection. DMSO
Feeding mice with *L. mesenteroides EH-1*. ICR mice were fed with live or heat (100 °C)-killed *L. mesenteroides EH-1* (8 × 10^8 CFU/50 µL) once a day for 2 days. Water (50 µL) was given as control. Two days after the *L. mesenteroides EH-1* feeding, cecum was homogenized in water (50 µg/500 µL), followed by a vortex mixing step, and stored at −80 °C until the detection of butyric acid by HPLC. Two weeks after daily feeding diabetic mice with *L. mesenteroides EH-1*, blood glucose was detected once a week, fasting blood was collected for the detection of insulin and IL-6 levels. Mice were sacrificed in a CO₂ chamber and the pancreases were collected for immunohistochemical analysis. For Ffar2 inhibition, diabetic mice were fed with *L. mesenteroides EH-1* (8 × 10^9 CFU/50 µL) once a day for 2 weeks. GLPG-0974 was given at 1 mL/kg body weight by gastric gavage just before *L. mesenteroides EH-1* administration and weekly. A vehicle was 0.1% DMSO in saline. Blood glucose was detected once a week. Two weeks after treatment, blood was collected for detection of insulin and IL-6 levels.

**Statistical analysis.** To determine significance between groups, comparisons were made using the two-tailed Student's t-test. Data are presented as mean values ± standard deviation (SD). The mean values ± SD for all figures with bar charts were shown in Supplementary Table 1. Statistical analyses were performed using GraphPad Prism 5 software. Unpaired Student's t-test was used to compare two groups. When appropriate, ANOVA was used and post hoc analysis was performed with Tukey's test to compare more than two groups. A p value < 0.05 was regarded as statistically significant.

**Results**

**Fermentation properties of *L. mesenteroides EH-1***. A single colony was isolated from a TSB agar plate spread with Mongolian curd cheese and evaluated by 16S rRNA sequencing. The 16S rRNA gene (Supplementary Fig. S1) of this colony shares 99% identity to that of *L. mesenteroides* ATCC 8293. This isolated strain was here named as *L. mesenteroides EH-1*. *L. mesenteroides* is a lactic acid bacterium that is currently used as a starter for kimchi and kefir²⁶. *L. mesenteroides EH-1* grew well at the temperatures of 25 °C and 37 °C, but not 4 °C (Supplementary Fig. S2). The growth of *L. mesenteroides EH-1* was unaffected by acidification of the media, as growth curves were similar from pH 3 to pH 7 (Supplementary Fig. S3). These results indicate that the strain of *L. mesenteroides EH-1* isolated from Mongolian curd cheese is stable at room temperature and tolerant of low pH. To examine the fermentative capabilities, *L. mesenteroides EH-1* was cultured in rich media in the presence of 2% glucose for 24 h. Rich media with glucose alone or *L. mesenteroides EH-1* alone served as controls. The media in the culture of *L. mesenteroides EH-1* with glucose turned yellow after incubation for 24 h, while the media in the other three conditions maintained their original colors (Fig. 1a). As shown in Fig. 1b,c, the OD_{562} and pH values of media with *L. mesenteroides EH-1* plus glucose demonstrated significant decreases compared to controls, indicating that *L. mesenteroides EH-1* has a capability of fermenting glucose. HPLC analysis was conducted to quantify the level of butyric acid in fermentation media of *L. mesenteroides EH-1*. Butyric acid is detectable in media from glucose fermentation of *L. mesenteroides EH-1*, but not media from controls (Fig. 1d). The different concentrations of butyric acid (0–100 mmol/L) were subjected to HPLC for establishment of a quantitative standard curve. As shown in Fig. 1e, glucose fermentation of *L. mesenteroides EH-1* for 24 h yielded approximately 1.6 mmol/L of butyric acid.

**Effects of butyric acid on insulin secretion from Min6 cells.** To investigate the effect of butyric acid on insulin secretion, Min6 cells were treated with butyric acid (100 mmol/L) for 24 h. Treatment of glucose or water served as positive and negative controls, respectively. Results in Fig. 2a showed that treatment of cells with butyric acid, like glucose, markedly elevated insulin levels in culture media of Min6 cells. To determine whether Ffar2 mediated the regulation of butyric acid on insulin secretion, cells were pre-treated with Ffar2 or scrambled siRNA before addition of water alone, glucose alone, or butyric acid plus glucose for 24 h. The Ffar2 siRNA induced a 74.46 ± 2.27% knockdown of Ffar2 gene (Supplementary Fig. S5). As shown in Fig. 2b, the knockdown of Ffar2 with its specific siRNA, but not scrambled siRNA, considerably blocked the effect of butyric acid on induction of insulin secretion. On the other hand, the Ffar2 knockdown had no influence on glucose-induced insulin secretion from Min6 cells. The results suggest that Ffar2 is essential for the action of butyric acid at induction of insulin secretion from Min6 cells.

**Involvement of Ffar2 in the effect of butyric acid on the levels of blood glucose, and insulin in type 1 diabetic mice.** To establish a type 1 diabetic mouse model, STZ was administered to ICR mice via IP injection. Compared to control mice, injection of STZ for two days led to higher levels of glucose and IL-6 in the blood. STZ-induced diabetic mice were injected intraperitoneally with butyric acid or water once a day for a week. Injection of butyric acid resulted in a remarkable decrease in fasting blood glucose (Fig. 3a) as well as IL-6 (Fig. 3c). Furthermore, the amount of insulin in the plasma of butyric acid–injected mice was higher than that of water-injected mice (Fig. 3b). To further confirm the essential role of Ffar2 in mediating the action of butyric acid in vivo, the STZ-induced diabetic mice were administered intragastrically with GLPG-0974, an Ffar2 antagonist, or DMSO control before injection of butyric acid or water. Administration of GLPG-0974, but not DMSO, counteracted the effect of butyric acid on the down-regulation of glucose and the up-regulation of insulin (Fig. 3d,e), although no effect on butyric acid-induced IL-6 reduction was observed (Fig. 3f). These results indicate that butyric acid may regulate the levels of glucose and insulin in blood of STZ-induced diabetic mice via binding to Ffar2.

**Production of butyric acid by *L. mesenteroides EH-1* in vivo.** To explore whether butyric acid can be produced in vivo by *L. mesenteroides EH-1*, ICR mice were fed with *L. mesenteroides EH-1* or water once a day for 2 days. Butyric acid in cecum homogenates was measured by HPLC. Butyric acid is detectable in the cecum
Figure 1. Glucose fermentation of *L. mesenteroides* EH-1. (a) *L. mesenteroides* EH-1 (LM) was incubated in rich media (M) with/without glucose (G) for 24 h. Rich media alone and rich media plus glucose without *L. mesenteroides* EH-1 were included as controls. Fermentation was detected by (b) OD_{565} and (c) pH value. (d) Butyric acid was measured by HPLC and (e) the concentrations of butyric acid were calculated from the height of butyric acid standard (STD) peaks. Data are the mean ± SD from 3 independent experiments. ***p < 0.001 vs M, ###p < 0.001 vs M + LM, and †††p < 0.001 vs M + G.

Figure 2. Effects of butyric acid on insulin secretion from Min6 cells. (a) Levels of insulin secretion from Min6 cells after treatments with water (H_{2}O), 100 µmol/L glucose (G), and 100 µmol/L butyric acid (BA) for 24 h were measured by ELISA. (b) Min6 cells pre-treated with Ffar2 or negative control siRNAs before incubation with water, glucose, and butyric acid for 24 h. The level (µg/L) of insulin was detected using a mouse insulin ELISA kit. Data are the mean ± SD from 3 independent experiments. **p < 0.01; ***p < 0.001 vs water treatment and #p < 0.05; ###p < 0.001 vs glucose treatment.
of *L. mesenteroides* EH-1-fed mice, but not in that of control mice fed with water (Fig. 4a,b). Approximately 0.7 mmol/L butyric acid was detected in the cecum, suggesting that *L. mesenteroides* EH-1 can produce butyric acid at a high concentration in a cecum microenvironment. To assess whether butyric acid‐producing *L. mesenteroides* EH-1 can lower the blood glucose in diabetes, STZ‐induced diabetic mice were fed with *L. mesenteroides* EH-1 or water. Feeding mice with *L. mesenteroides* EH-1 once a day for 2 weeks substantially reduced the levels of glucose (Fig. 4c) and IL‐6 in the blood (Fig. 4e) and increased the amounts of insulin (Fig. 4d) in the blood and pancreas (Fig. 4d, inserted panels). These results demonstrate the probiotic activity of *L. mesenteroides* EH-1 in regulating the levels of glucose and IL‐6 in STZ‐induced type 1 diabetic mice. To validate the contribution of Ffar2 to the effect of butyric acid‐producing *L. mesenteroides* EH-1 on lowering blood glucose, GLPG-0974 was given to STZ‐induced diabetic mice to antagonize the Ffar2 before feeding mice with *L. mesenteroides* EH-1. Compared to mice treated with DMSO control, mice given GLPG-0974 displayed no difference in glucose (Fig. 5a), IL‐6 (Fig. 5c), as well as insulin (Fig. 5c) in blood. Taken together, the data in Figs. 3 and 5 strongly suggest that Ffar2 mediates the signaling of butyric acid produced by *L. mesenteroides* EH-1 to diminish the elevated glucose in STZ‐induced diabetic mice.

**Discussion**

Probiotic *L. mesenteroides* is currently used for food fermentation. An addition of *L. mesenteroides* to cabbage fermentation ensured that texture and flavor quality were retained, while providing a 50% reduction in sodium chloride67. In addition, this probiotic bacterium has demonstrated a number of beneficial effects, including inhibition of three fish pathogens68, suppression of low‐pathogenic avian influenza (H9N2) virus in chickens69, and reduction of *Streptococcus thermophilus* induced IL‐12 and interferon (IFN)‐γ production in human peripheral blood mononuclear cells69.
Previous studies have identified potential antidiabetic effects of other commensal bacteria. Treatment with *Lactobacillus casei* CCFM419 improved impaired pancreatic function and attenuated type 2 diabetes in a mouse model. Carrot juice fermentation of *Lactobacillus plantarum* NCU116 reduced pancreatic injuries. Oral administration of *Lactobacillus rhamnosus* CCFM0528 improved glucose intolerance by protecting islet cells. Our current study demonstrated for the first time that *L. mesenteroides* EH-1 exerts a probiotic activity that lowers blood glucose in the STZ-induced type 1 diabetic mouse model. To validate the essential role of *L. mesenteroides* EH-1 in regulation of glucose and insulin levels in diabetic mice, we fed STZ-induced type 1 diabetic mice with heat-killed *L. mesenteroides* EH-1. As shown in Supplementary Fig. S7, the killed *L. mesenteroides* EH-1 bacteria lost their capacity to lower glucose and increase insulin, indicating that *L. mesenteroides* EH-1, not endogenous gut microbes indirectly affected by probiotic *L. mesenteroides* EH-1, exerted the anti-diabetic properties. Future works will use germ-free mice or mice with the gut microbiome depleted by antibiotics to study the effects of probiotic bacteria and endogenous gut microbes on the regulation of blood glucose and insulin.

Ffar2 is highly expressed on pancreatic islets and up-regulated in islets during pregnancy. The potency rank order of SCFAs for Ffar2 is acetate (C2) ~ propionate (C3) > butyrate (C4) > valerate (C5) > formate. Moreover, Ffar2 is a novel effector of glucose homeostasis in part due to its direct effect on insulin secretion and beta cell proliferation. In agreement with previous studies, our data revealed that treatment of Min6 cells with 100 mmol/L butyric acid down regulated insulin receptor (IR), and IR substrate 1 (IRS-1) expression involved in insulin signaling in the mouse liver. Application of butyrate also associated with decreased insulin receptor beta subunit (IR-beta)
expression of in hepatic tissue. Thus, administration of butyric acid into mice may enhance the insulin secretion and prolong the insulin clearance in mice.

Butyric acid is one of Ffar2’s known agonists. Previous studies have shown that Ffar2 agonism can trigger an increase in intracellular inositol triphosphate and Ca^{2+} levels, and potentiate insulin secretion. In addition, sodium butyrate treatment improved glucose homeostasis and reduced beta cell apoptosis in diabetic rats. Knockdown of Ffar2 significantly reduced butyric acid-induced insulin secretion from Min6 cells, clearly illustrating that Ffar2 signaling is an important effector of insulin secretion induced by butyric acid. Previous studies have demonstrated a strong reduction in plasma glucose by feeding with ketogenic diets. Both propionic acid and butyric acid are ketogenic substrates and can bind to Ffar2. The effects of propionic acid and butyric acid on insulin secretion from Min6 cells were compared side-by-side. Propionic acid induced detectable amounts of insulin secreted from Min6 cells although it is less effective than butyric acid (Supplementary Fig. S6). Knockdown of Ffar2 considerably lowered propionic acid-induced insulin secretion from Min6 cells. Butyric acid, but not propionic acid, is a potent inhibitor of histone deacetylases (HDAC). Previous studies demonstrated that butyrate can improve insulin sensitivity via HDAC inhibition. Butyrate as a dietary supplement can prevent high fat diet-induced insulin resistance in mice by promotion of energy expenditure and induction of mitochondria function. Fasting insulin was significantly lower in the butyrate-treated high-fat diet mice. The signal of phosphorylation of IRS-1 in the skeletal muscle was increased in butyrate-treated mice, suggesting a molecular mechanism of insulin sensitization. Our data revealed that butyric acid can mediate Ffar2 to increase blood insulin in STZ-induced type 1 diabetic mice. Thus, it is worth investigating whether butyric acid controls the activity of Ffar2 or HDAC to regulate the insulin secretion in different types of diabetes.

Loss of Ffar2 in mice increases the risk of diabetic status, since Ffar2 knockout (KO) mice exhibit fasting hyperglycemia, reduced insulin levels, and glucose intolerance, despite exhibiting normal insulin sensitivity. Exposing Ffar2 KO mice to a high fat diet resulted in a decrease in both islet number and size, leading to reduced beta cell mass and total pancreatic insulin content. A butyrate-enriched diet could partially protect Ffar2 KO mice in a non-obese diabetic background from type 1 diabetic islet inflammation. As shown in Fig. 3, a single dose of GLPG-0974, a Ffar2 antagonist, suppressed butyric acid-induced increase of fasting blood insulin and decreased fasting blood glucose, confirming the indispensable role of Ffar2 in the action of butyric acid in vivo. Furthermore, Ffar2 activation on intestinal enteroendocrine cells induces the production of glucagon-like peptide...
(GLP)-1 which is anorexigenic and stimulates insulin secretion. Future work is required to determine if GLP-1 mediates the Ffar2-butyric acid induced insulin secretion in the STZ-induced type 1 diabetic mouse model.

STZ specifically damages pancreatic islet beta cells. The damaged beta cells passively relate to high mobility group box 1 (HMGB1) release. In parallel, the inflammatory cells infiltrated pancreatic islets, such as macrophages, dendritic cells and T cells. The differentiation of Naive CD4+ T cells into effector T helper (Th)1 and/or Th17 cells based on current cytokine microenvironment. In addition, the released HMGB1 targets macrophage or dendritic cells via the corresponding surface receptor(s), which induces a cascade signal that activates the nuclear factor κB (NF-κB) pathway. This results in the activation of Th17 cells, therefore leading to the production of proinflammatory cytokines IL-1β, TNF-α, and IL-6. It has been reported that IL-6 induces beta cell apoptosis via signal transducer and activator of transcription (STAT)-3-mediated the production of nitric oxide. The dysregulation of proinflammatory cytokines and Th1/Th2/Th17 further accelerate an inflammation of islets and destruction of beta cells, and leads to type 1 diabetes. Sodium butyrate can heal the balance of Th1/Th2 and block Th17 cells.

In this study, we found that the acute blocking of Ffar2 by GLPG-0974 at a single dose for 24 h showed a minor effect of Ffar2-butyric acid signaling on IL-6 reduction (Fig. 3f), whereas the long term blocking of Ffar2 using multiple doses of GLPG-0974 for two weeks resulted in a significant the suppression effect on the L. mesenteroides EH-1-reduced blood IL-6 level in type 1 diabetic mice (Fig. 3c). Although other Ffar2 antagonists from azetidine derivatives at different doses can be used to completely block Ffar2 in vivo, it has been documented that sodium butyrate, as a direct HMGB1 antagonist, could down-regulate the expression of HMGB1 and mediate the balance of Th1/Th2/Th17 paradigm, thus attenuating type 1 diabetes. Thus, it is possible that butyric acid reduced IL-6 production by directly down-regulating HMGB1 expression and bypassing Ffar2.

STZ does not influence the function of pancreatic beta cells of humans when used in the treatment of islet-cell carcinomas. Insulin can be detected in STZ-injected mice (Fig. 4d), indicating that STZ injection results in incomplete damage to the pancreas in mice. It has been shown that Ffar2 directly mediates both the stimulatory effects of sodium acetate and propionate on insulin secretion and their protection against islet apoptosis. In addition to butyric acid, other SCFAs in the fermentation media of L. mesenteroides EH-1 will be measured in the future. One of limitations of using SCFAs as therapeutics includes their typically short half-life, with clearance from plasma occurring within a few hours. A high concentration (about 0.7 mmol/L) of butyric acid was detected in the cecum of mice fed with L. mesenteroides EH-1 (Fig. 4a), demonstrating that L. mesenteroides EH-1 is a potent strain for producing butyric acid. Data in literature demonstrated that high-fat-fed mice treated with butyrate showed enhancement of the insulin secretion, which was related to a substantial reduction in lipid accumulation within the pancreas. Furthermore, butyrate can prevent type 1 diabetes in non-obese diabetic (NOD) mice. Results from previous studies above supported that butyric acid has a beneficial effect on prevention of type 1 diabetes. Future works will test the anti-diabetic activity of probiotic L. mesenteroides EH-1 using NOD mice or high-fat diet-induced type 1 diabetes.

In summary, our study characterized a new probiotic bacterial strain, L. mesenteroides EH-1, which was originally isolated from Mongolian curd cheese. L. mesenteroides EH-1 produces high concentrations of butyric acid which can activate Ffar2 to raise insulin levels but mitigate glucose amounts in the blood of type 1 diabetic mice.

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References

1. Warren, S. & Root, H. F. The pathology of diabetes, with special reference to pancreatic regeneration. Am. J. Pathol. 1, 415 (1925).
2. Gepts, W. Pathologic anatomy of the pancreas in juvenile diabetes mellitus. Diabetes 14, 619–633 (1965).
3. Junker, K., Egeberg, J., Kromann, H. & Nerup, J. An autopsy study of the islets of Langerhans in acute-onset juvenile diabetes mellitus. Acta Path. Microbiol. Scand. Sect. A 85, 699–706 (1977).
4. Pipeleers, D. & Ling, Z. Pancreatic beta cells in insulin‐dependent diabetes. Diabetologia 54, 2325–2340 (2011).
5. Calcium, R., et al. Effects of sodium acetate and propionate on insulin secretion and their protection against islet-cell carcinomas. Immunity 65, 633–664 (2016).
6. Eizirik, D. L., et al. The role of short-chain fatty acids in the interplay between diet, gut microbiota, and host energy metabolism. J. Lipid Res. 54, 2325–2340 (2013).
7. Al-Salami, H. & Nash, A. A. Effect of probiotics on glucose metabolism in patients with type 2 diabetes mellitus: a meta-analysis of randomized controlled trials. Medicine 52, 28–34 (2016).
8. Morrison, D. J. & Presto, T. Development of short chain fatty acids by the gut microbiota and their impact on human metabolism. Gut microbes 7, 189–200 (2016).
9. Priyadarshini, M., Wicksteed, B., Schiltz, G. E., Gilchrist, A. & Layden, B. T. SCFA receptors in pancreatic β-cells: novel diabetes targets? Trends Endocrinol. Metab. 27, 563–566 (2014).
10. Bhutia, Y. D. & Ganapathy, V. Short, but smart: SCFAs train T cells in the gut to fight autoimmunity in the brain. Immunity 43, 629–631 (2015).
11. den Besten, G. et al. The role of short-chain fatty acids in the interplay between diet, gut microbiota, and host energy metabolism. J. Lipid Res. 54, 2325–2340 (2013).
48. Leonard, J. N. & Hakak, Y. (Google Patents, 2010).
49. Layden, B. T.
50. McNelis, J. C.
51. Matis, G.
52. Meehan, A. E.
53. Mariño, E.
54. Jin, C. J., Sellmann, C., Engstler, A. J., Ziegenhardt, D. & Bergheim, I. Supplementation of sodium butyrate protects mice from the onset of type 1 diabetes in young children. *Diabetologia* 57, 1569–1577 (2014).
55. Endesfelder, D. et al. Towards a functional hypothesis relating anti-islet cell autoimmunity to the dietary impact on microbial communities and butyrate production. *Microbiome* 4, 17 (2016).
56. Kleytmans, J., Van Belkum, A. & Verbrugh, H. Nasal carriage of *Staphylococcus aureus*: epidemiology, underlying mechanisms, and associated risks. *Clin. Microbiol. Rev.* 10, 505–520 (1997).
57. Round, J. L. et al. The Toll-like receptor 2 pathway establishes colonization by a commensal of the human microbiota. *Science*, 1206095 (2011).
58. Furman, B. L. Streptozotocin-induced diabetic models in mice and rats. *Curr. Protoc. Pharmacol.* 70, S.4.7, 41–45.47. 20 (2015).
59. King, A. J. The use of animal models in diabetes research. *British journal of pharmacology* 166, 877–894 (2012).
60. Somboonwong, J., Traisaeng, S. & Saguanrungsirikul, S. Moderate-intensity exercise training elevates serum and pancreatic zinc levels and pancreatic ZnT8 expression in streptozotocin-induced diabetic rats. *Life Sci.* 139, 46–51 (2015).
61. Akba, Y. et al. FFA2 activation combined with ulroglicenic COX inhibition induces duodenal mucosal injury via the 5-HT pathway in rats. *Am. J. Physiol-Gastrl.* 313, G117–G128 (2017).
62. Kim, J. E. et al. Enhancing acid tolerance of *Leuconostoc mesenteroides* with gluthathione. *Biotechnol. Lett.* 34, 683–687 (2012).
63. Johanningsmeier, S., McFeeters, R. F., Fleming, H. P. & Thompson, R. L. Effects of *Leuconostoc mesenteroides* starter culture on metabolite production. *Exp. Patho.* 4034–4043 (2007).
64. Jakobsdottir, G., Xu, J., Molin, G., Ahnre, S. & Nyman, M. High-fat diet reduces the formation of butyrate, but increases succinate, inflammation, liver fat and cholesterol in rats, while dietary fibre counteracts these effects. *PloS one* 8, e80476 (2013).
65. Yadav, H., Lee, J.-H., Lloyd, J., Walter, P. & Rane, S. G. Beneficial metabolic effects of a probiotic via butyrate induced GLP-1 secretion. *J. Biol. Chem., jbc.* M113, 452516 (2013).
66. de Goffau, M. A. C. et al. Aberrant gut microbiota composition at the onset of type 1 diabetes in young children. *Diabetologia* 57, 1569–1577 (2014).
67. Li, C. et al. Carrot juice fermented with *Lactobacillus plantarum* NCU116 ameliorates type 2 diabetes in rats. *J. Agr. Food Chem.* 62, 11884–11891 (2014).
68. Chen, P. et al. Oral administration of *Lactobacillus rhamnosus* CCFM4191 on insulin resistance and gut microbiota in type 2 diabetic mice. *Benef. Microbes* 8, 421–432 (2017).
69. Li, C. et al. Carrot juice fermented with *Lactobacillus plantarum* NCU116 ameliorates type 2 diabetes in rats. *J. Agr. Food Chem.* 62, 11884–11891 (2014).
70. Chen, P. et al. Oral administration of *Lactobacillus rhamnosus* CCFM4528 improves glucose tolerance and cytokine secretion in high-fat-fed, streptozotocin-induced type 2 diabetic mice. *J. Funct. Foods.* 10, 318–326 (2014).
71. Kozakova, H., Schwarzer, M., Srtukova, D., Hudovici, T. & Cukrowska, B. Colonisation of germ-free mice with probiotic lactobacilli mitigates allergic sensitisation in murine model of birch pollen allergy. *Clinical and translational allergy* 4, P26 (2014).
72. Zarrinpar, A. et al. Antibiotic-induced microbiome depletion alters metabolic homeostasis by affecting gut signaling and colonic metabolism. *Nature communications* 9, 1–13 (2018).
73. Regel, J. B. et al. Probing cell type–specific functions of G i in vivo identifies GPCR regulators of insulin secretion. *J. Clin. Invest.* 117, 4034–4043 (2007).
74. Ahren, B. Islet G protein-coupled receptors as potential targets for treatment of type 2 diabetes. *Nat. Rev. Drug Discov.* 8, 369 (2009).
75. Leonard, N. & Hakala, Y. (Google Patents, 2010).
76. Layden, B. T. et al. Regulation of pancreatic islet gene expression in mouse islets by pregnancy. *J. Endocrinol.* 207, 265–279 (2010).
77. Ulven, T. Short-chain free fatty acid receptors FFA2/GPR43 and FFA3/GPR41 as new potential therapeutic targets. *Front. Endocrinol.* 3, 111 (2012).
78. Fuller, M. et al. The short-chain fatty acid receptor, FFA2, contributes to gestational glucose homeostasis. *Am. J. Physiol-Endoc.* M. 309, E840–E851 (2015).
79. McNelis, J. C. et al. GPR43 potentiates beta cell function in obesity. *Diabetes*, db141938 (2015).
80. Buckworth, J. W. C., Bennett, R. G. & Harnel, F. G. Insulin degradation: progress and potential. *Endocrine reviews* 19, 608–624 (1998).
81. Jin, C., Sellmann, C., Engstler, A. J., Ziegenhardt, D. & Bergheim, I. Supplementation of sodium butyrate protects mice from the development of non-alcoholic steatohepatitis (NASHi). *British journal of Nutrition* 114, 1745–1755 (2015).
82. Raso, G. M. et al. Effects of sodium butyrate and its synthetic amide derivative on liver inflammation and glucose tolerance in an animal model of steatosis induced by high fat diet. *PloS one* 8 (2013).
83. Mattis, G. et al. Effects of oral butyrate application on insulin signalling in various tissues of chickens. *Domestic animal endocrinology* 50, 26–31 (2015).
84. Boros, L. G., Collins, T. G. & Somlyay, G. What to eat or not to eat—or what not to eat—is still the question. *Neuro-oncology* 19, 595–596 (2017).
85. Davie, J. R. Inhibition of histone deacetylase activity by butyrate. *The journal of nutrition* 133, 2485S–2493S (2003).
86. Martin, E. et al. Gut microbial metabolites limit the frequency of autoimmune T cells and protect against type 1 diabetes. *Nat. Immunol.* 18, 552 (2017).
87. Bindels, L. B., Dewulf, E. M. & Delzenne, N. M. GPR43/FFA2: physiopathological relevance and therapeutic prospects. *Trends Pharmacol. Sci.* 34, 226–252 (2013).
88. Han, J. et al. Extracellular high-mobility group box 1 acts as an innate immune mediator to enhance autoimmune progression and diabetes onset in NOD mice. *Diabetes* 57, 2118–2127 (2008).
89. Zhang, S., Zhang, L., Yang, P., Gong, F. & Wang, C.-Y. HMGB1, an innate alarmin, in the pathogenesis of type 1 diabetes. *Int. J. Clin. Exp. Patho.* 3, 24 (2010).
90. Abdel-Moneim, A., Bakery, H. H. & Allam, G. The potential pathogenic role of IL-17/Th17 cells in both type 1 and type 2 diabetes mellitus. *Biomed. Pharmacother.* 101, 287–292 (2018).
64. Oh, Y. S., Lee, Y. J., Park, E. Y. & Jun, H. S. Interleukin-6 treatment induces beta-cell apoptosis via STAT-3-mediated nitric oxide production. Diabetes Metab. Res. Rev. 27, 813–819 (2011).
65. Guo, Y. et al. Sodium butyrate ameliorates streptozotocin-induced type 1 diabetes in mice by inhibiting the HMGB1 expression. Front. Endocrinol. 9, 630 (2018).
66. Hansen, A. H. et al. Development and characterization of a fluorescent tracer for the free fatty acid receptor 2 (FFA2/GPR43). J. Med. Chem. 60, 5638–5645 (2017).
67. Eleazu, C. O., Eleazu, K. C., Chukwuma, S. & Essien, U. N. Review of the mechanism of cell death resulting from streptozotocin challenge in experimental animals, its practical use and potential risk to humans. J. Diabetes Metab. Disord. 12, 60 (2013).
68. Pingitore, A. et al. Short chain fatty acids stimulate insulin secretion and reduce apoptosis in mouse and human islets in vitro: Role of free fatty acid receptor 2. Diabetes Obes. Metab. 21, 330–339 (2019).
69. Pace, B. S. et al. Short-chain fatty acid derivatives induce fetal globin expression and erythropoiesis in vivo. Blood 100, 4640–4648 (2002).
70. Matheus, V., Monteiro, L., Oliveira, R., Maschio, D. & Collares-Buzato, C. Butyrate reduces high-fat diet-induced metabolic alterations, hepatic steatosis and pancreatic beta cell and intestinal barrier dysfunctions in prediabetic mice. Experimental Biology and Medicine 242, 1214–1226 (2017).

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Author contributions
S.T. and C.M.H. conceived and designed experiments, acquired and analyzed data and wrote the manuscript. A.B. and B.C. were in charge of bacterial isolation and characterization from Mongolian curd cheese, Y.F. acquired HPLC analysis and T.H.C. analyzed data and review manuscript. D.R.H. reviewed the manuscript. All authors approved the final version of the manuscript. C.M.H. is the guarantor of this work.

Competing interests
The authors declare no competing interests.

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