Evidence continues to emerge detailing the myriad of ways the gut microbiota influences host energy homeostasis. Among the potential mechanisms, short chain fatty acids (SCFAs), the byproducts of microbial fermentation of dietary fibers, exhibit correlative beneficial metabolic effects in humans and rodents, including improvements in glucose homeostasis. The underlying mechanisms, however, remain elusive. We here report that one of the main bacterially produced SCFAs, propionate, activates ileal mucosal free fatty acid receptor 2 to trigger a negative feedback pathway to lower hepatic glucose production in healthy rats in vivo. We further demonstrate that an ileal glucagon-like peptide-1 receptor-dependent neuronal network is necessary for ileal propionate and long chain fatty acid sensing to regulate glucose homeostasis. These findings highlight the potential to manipulate fatty acid sensing machinery in the ileum to regulate glucose homeostasis.

Obesity and diabetes associate with shifts in the gut microbiome. Among the proposed mechanisms underlying the role of the microbiota in host energy homeostasis, production of short chain fatty acids (SCFAs) from non-digestible carbohydrates is hypothesized to be a main contributor (1). Alterations in SCFA production, either via prebiotic supplementation or surgical alteration, improve metabolic parameters (2–5). The three main SCFAs, acetate, propionate, and butyrate, differentially alter host metabolism (2, 3), and it appears that propionate exerts the most beneficial metabolic outcome as body weight changes via gastric bypass associate with rises in propionate (4), mice fed a high-fat diet supplemented with propionate improves glucose tolerance (5), and delivery of propionate to the colon of obese humans reduces weight gain (6).

SCFAs act on free fatty acid receptor-2 (FFAR2) and -3 (FFAR3), formerly termed GPR43 and GPR41, respectively (7, 8), which are expressed in intestinal epithelial cells (9, 10). Both receptors are localized in enteroendocrine cells (11) and SCFA-induced glucagon-like peptide-1 (GLP-1) release is impaired in primary intestinal cultures from FFAR2−/− and FFAR3−/− mice (12), whereas intestinal infusion of propionate fails to induce gut peptide release in FFAR2−/−, but not FFAR3−/− mice (13). Of note, these results are associative indications of intestinal FFAR2 regulating blood glucose levels via GLP-1, but to date, no one has addressed a direct causative role of intestinal FFAR2 activation in the regulation of glucose homeostasis in vivo.

To this end, we have characterized the role of the proximal small intestine in the regulation of glucose homeostasis, whereby duodenum and jejunum sensing mechanisms lower hepatic glucose production (GP) via a gut-brain-liver neuronal network involving gut peptide signaling (14–18). Despite a contribution in the control of food intake via activation of the ileal brake (19), much less is known about the glucoregulatory role of the ileal sensing (20). Given that FFAR2 is highly expressed in the ileum (9), and that conditions in the ileum are more favorable for microbial growth and SCFA production than the proximal small intestine (21, 22), we postulated that propionate triggers an ileal neuronal-dependent negative feedback pathway that requires ileal FFAR2 and is mediated via GLP-1 receptor (GLP-1R) signaling to lower GP and maintain whole body glucose homeostasis. Furthermore, ileal long chain fatty acids (LCFAs) stimulate GLP-1 secretion (23, 24) and vagal afferent firing (25). Thus, to further characterize the general ileal fatty acid-sensing pathways, we additionally investigated, for the first time to our knowledge, whether LCFAs sensing in the ileum regulates glucose homeostasis and whether a similar GLP-1R neuronal network is required.

**Experimental Procedures**

*Animals*—Male Sprague-Dawley rats (250–300 g) were obtained from Charles River Laboratories (Montreal, QC) and were housed in individual cages maintained on a 12-h light/dark cycle. Rats had access to chow and water *ad libitum*, and...
were given 7 days to acclimatize before the designated surgeries were performed. Animals were randomly designated into experimental groups. The protocols were approved by the Institutional Animal Care Committee of UHN.

**Ileal and Intravenous Cannulations**—A gut catheter was inserted 2-cm proximal to the cecum to target the ileum and allow for infusion of compounds and blood sampling. Rats were given 3–4 days to recover from the surgery before the infusion studies.

**Viral Injection and Surgery**—For experiments involving ileal FFAR2 knockout, we utilized established protocols that target infection to a selective segment of the small intestine (16, 17, 26). Lentivirus (LV) expressing FFAR2 shRNA (1.0 × 10^6 infectious units of virus; Santa Cruz Biotechnology Inc., Dallas, TX) or mismatch (1.0 × 10^6 infectious units of virus; Santa Cruz Biotechnology Inc.) was diluted 1/10 into 0.2 ml of saline and injected into the ileal lumen and left for 20 min to optimize infection. An ileal catheter was subsequently inserted through the injection site, followed by vascular cannulation. Animals were allowed to recover for 3 days before the infusion studies.

**Pancreatic (Basal Insulin)-Euglycemic Clamp with Steady-state Glucose Kinetics Assessment**—This procedure was performed as described (15, 18). The night before the infusion-clamp studies, rats were fasted for 4–6 h. The pancreatic clamp technique (Fig. 1A) was performed in conjunction with tracer glucose-dilution methodology, allowing for the simultaneous calculation of GP and glucose uptake during intralreal fatty acid infusions. The experiments lasted a total of 200 min, with a continuous infusion of [3-3H]glucose (40 µCi bolus; 0.4 µCi min⁻¹; PerkinElmer Life Sciences) initiated at the onset of the experiment (t = 0 min) and maintained throughout (t = 200 min). After reaching steady-state of the [3-3H]glucose infusion, the pancreatic clamp was initiated at t = 90 min (insulin at 1.2 milliunits kg⁻¹ min⁻¹ and somatostatin at 3 µg kg⁻¹ min⁻¹). Plasma samples were taken every 10 min and exogenous 25% glucose solution was infused accordingly to maintain euglycemia. The following treatments were given into the ileum (0.01 ml min⁻¹ for 50 min): (a) saline, (b) 1% bile acid (phosphate-buffered saline (PBS) with lecithin and sodium taurocholate), (c) propionate (Prop; 5 mM; 4.8 µg min⁻¹), (d) 5% Intralipid (IL; 20% Intralipid diluted in saline; Baxter Corporation), (e) oleic acid (OA; 200 µm; 0.564 mg min⁻¹), (f) linoleic acid (LA; 200 µm; 0.56 mg min⁻¹), (g) triacis-C (0.48 pmol min⁻¹), (h) exendin-9 (0.15 µg min⁻¹; Tocris Bioscience, Ellisville, MO), and (i) tetracaine (0.01 mg min⁻¹; Sigma; this dose selected based on a previous study that evaluated upper small intestinal fatty acid sensing (18)). Sodium propionate (>99% purity; Sigma) powder was dissolved in saline to make 5 mM propionate solution. This dose of propionate was chosen based on the fact that previous studies have mostly examined colonic SCFA mechanisms (13), and as colonic SCFA levels are likely to be supraphysiological as compared with the ileum (27), we chose a lower dose of SCFA (5 mM) than the concentration that was used in the colon. In addition, 5 mM SCFA has been documented to stimulate GLP-1 secretion in vitro (13) and more importantly, this dose of propionate is comparable with ileal propionate levels reported in humans (28). Intralipid was infused at one-fourth of the dose that was used in duodenal studies (18) because the amount of Intralipid detected in the ileum is lower than the duodenum following intragastric administration (29). Oleic or linoleic acids (99% purity; Sigma) were emulsified in PBS, 1% sodium taurocholate (Sigma), and 0.1% lecithin (L-α-phosphatidylcholine; Sigma) to make 200 mM solutions as described (30). A concentration of 200 mM was chosen for oleic or linoleic acids because this concentration is comparable with what was reported to stimulate gut peptide release when infused into the ileum (23, 31).

**Protein Extraction and Western Blotting**—Tissues were lysed on ice with a handheld homogenizer in a buffer containing: 50 mM Tris–hydrochloric acid (pH 7.5), 1 mM EGTA, 1 mM EDTA, 1% (w/v) Nonidet P-40, 1 mM sodium orthovanadate, 50 mM sodium fluoride, 5 mM sodium pyrophosphate, 0.27 M sucrose, 1 mM dithiothreitol (DTT) and protease inhibitor mixture (Roche Diagnostics, Laval, QC, Canada), and concentrations were determined with the Pierce 660 nm protein assay (Thermo Scientific, Rockford, IL). 30 µg of tissue lysates were subjected to electrophoresis in a mini–protein precast gel (4–15%; Bio-Rad) and transferred to nitrocellulose membranes. The membranes were incubated for 30 min with TBS-T containing 5% BSA. The membranes were then incubated with the indicated primary antibodies: anti-FFAR2 (1:200; sc-28424, Santa Cruz Biotechnology Inc.) or anti-β-actin (1:30,000, Sigma, catalog number A1978). The blots were then incubated with appropriate secondary antibodies (diluted 1:3000 for FFAR2 and 1:40,000 for actin) for 1 h in 5% skim milk at room temperature. After repeating the washing steps, the signal was detected with the enhanced chemiluminescence reagent. Western blots were developed with the Microchemi 2.0 DNR Bio-imaging system, and quantified protein levels with GelQuan software (DNR Bio-imaging system) and normalized to β-actin.

**Biochemical Analysis**—Plasma glucose levels were measured using the glucose oxidase method (GM9 Glucose Analyzer, Analox Instruments). Plasma insulin levels were determined by radioimmunoassay (Linco Research, St. Charles, MO). Plasma FFAs were measured by an enzymatic assay (Wako Pure Chemical Industries, Osaka, Japan).

**Statistical Analysis**—Results were analyzed using unpaired Student’s t test, or ANOVA (followed by Tukey’s post hoc test), using GraphPad Prism 5.0d software. Differences were considered significant at p < 0.05. Data are presented as mean ± S.E. The time period of 60–90 min was averaged for the basal condition, and the time period of 180–200 min was averaged for the clamp condition.
**Ileal Fatty Acid Sensing and Glucose Regulation**

![Diagram](https://via.placeholder.com/150)

**FIGURE 1. The effect of intraluminal propionate infusion on glucose kinetics.** A, schematic representation of the working hypothesis (left) and experimental procedure and pancreatic (basal insulin) euglycemic clamp protocol (right). SRIF, somatostatin. B–E, the glucose infusion rate (B), rate of GP (C), % suppression of GP (D), and glucose uptake (E) during the pancreatic clamp in rats with intraduodenal saline (Sal; n = 6) or propionate (Prop; n = 7) infusions, or intravenous (i.v.; n = 6) propionate infusion. Values are shown as mean ± S.E. **, p < 0.01; ***, p < 0.001, versus all other groups as determined by ANOVA with Tukey’s post hoc test.

**Results**

**Intraileal Propionate Infusion Lowers GP via Preabsorptive Mechanisms**—We first examined how whole body glucose metabolism changes in response to an acute rise of propionate in the ileum. Propionate was infused directly into the ileal lumen of awake, unrestrained healthy rats at a dose (5 mM) comparable with previous studies (13) and consistent with ileal propionate levels reported in humans (28). Insulin was maintained at plasma basal levels, which was achieved by the euglycemic–pancreatic clamp (Fig. 1A, Table 1). When propionate was infused into the ileal lumen for only 50 min, the exogenous glucose infusion rate required to maintain euglycemia was elevated compared with intraduodenal saline-infused rats (Fig. 1B), due to an inhibition of GP (Fig. 1, C and D) and not to an increase in the rate of glucose uptake, as peripheral glucose uptake did not change under these experimental conditions (Fig. 1E). Plasma FFA levels did not rise following intraluminal propionate infusions (Table 1), suggesting that the effect of propionate was preabsorptive. Consistently, intravenous infusion of propionate administered at the same dose and duration as intraileal delivery (i.e. assuming 100% leakage into the circulation following ileal infusion) did not alter the glucose infusion rate, the rate of GP, or glucose uptake (Fig. 1, B–E). Therefore, although the physiological relevance of ileal SCFA sensing has yet to be tested, these data demonstrate that ileal propionate sensing triggers a negative feedback pathway to lower GP via preabsorptive mechanisms.

**FFAR2 in the Ileum Is Necessary for Propionate Sensing**—Given that propionate is a potent ligand of FFAR2 (7) and that FFAR2 is highly enriched in the ileal mucosa (9), we examined whether FFAR2 is required for ileal propionate sensing (Fig. 2A). First, Western blot analysis revealed that FFAR2 (37 kDa) was expressed in the ileal mucosa of rats, using the respective testes as a negative control (32) (Fig. 2B) and confirming previous findings obtained in humans and rodents (7, 8, 33), although the relative expression of FFAR2 in the ileal enteroendocrine cells versus other mucosal cells remains to be determined. To investigate the necessity of ileal FFAR2 in glucose control, a previously established intestinal targeted viral approach was employed to knockdown FFAR2 protein selectively in the ileum in vivo (16, 17, 26). Three days before the infusion experiments and prior to ileal cannulations, a lentiviral expressing either mismatched shRNA (LV-MM) or FFAR2 shRNA (LV-FFAR2) was injected into the ileal lumen. Western blot analysis validated the knockdown of FFAR2 protein in the ileal mucosa of LV-FFAR2 versus rats injected with LV-MM (~51% reduction; Fig. 2C), with no changes in FFAR2 protein expression in the jejunal mucosa (Fig. 2D) or liver (Fig. 2E) of virus-injected rats, confirming the infection was localized to the ileum.

Rats injected with LV-MM or LV-FFAR2 had similar glucose kinetics in comparison to non-viral rats following a 50-min clamp.

**TABLE 1**

|              | Vehicle | Propionate | Oleic acid | Linoleic acid |
|--------------|---------|------------|------------|---------------|
| **Basal**    |         |            |            |               |
| Insulin (ng/ml) | 0.9 ± 0.1 | 0.8 ± 0.2 | 0.9 ± 0.1 | 0.9 ± 0.1      |
| Glucose (mM)       | 8.2 ± 0.4 | 6.2 ± 0.2 | 6.2 ± 0.2 | 7.0 ± 0.3      |
| Free fatty acids (mM) | 0.8 ± 0.1 | 0.8 ± 0.2 | 0.9 ± 0.1 | 0.8 ± 0.1      |
| **Clamp**       |         |            |            |               |
| Insulin (ng/ml) | 0.8 ± 0.2 | 0.8 ± 0.1 | 0.8 ± 0.1 | 0.9 ± 0.1      |
| Glucose (mM)       | 7.5 ± 0.1 | 6.6 ± 0.2 | 6.6 ± 0.2 | 7.5 ± 0.3      |
| Free fatty acids (mM) | 0.7 ± 0.1 | 0.8 ± 0.1 | 0.8 ± 0.1 | 0.7 ± 0.1      |
intraileal saline infusion (Fig. 3, A–D). Intraileal infusion of propionate for 50 min increased the glucose infusion rate (Fig. 3A) and lowered GP (Fig. 3, B and C), but did not alter the rate of glucose utilization (Fig. 3D) in LV-MM-injected control rats compared with intraileal saline infusion. Importantly, no difference in the glucose infusion rate, GP, or glucose uptake in LV-FFAR2 shRNA-injected rats infused with intraileal propionate was detected compared with intraileal saline-infused rats (Fig. 3, A–D).

To further establish site specificity and rule out the possibility that our ileal propionate infusion is activating colonic sensing mechanisms to lower GP, we performed an identical experiment to that above, except that LV-FFAR2 shRNA was injected into the proximal colon 3 days before clamp studies. Intraileal propionate infusion still resulted in a high glucose infusion rate (4.7 ± 0.4 mg kg⁻¹ min⁻¹) needed to maintain euglycemia, and lowered GP from basal (11.7 ± 0.3 versus 5.3 ± 0.3 mg kg⁻¹ min⁻¹; basal versus clamp) in rats with proximal colon LV-FFAR2 shRNA injection (n = 5). Taken together, intraileal propionate sensing activates ileal mucosa FFAR2 to lower GP in vivo.

An Ileal GLP-1 Receptor-dependent Neuronal Network Is Necessary for Propionate Sensing—Given that intestinal FFAR2 activation increases GLP-1 secretion (12, 13), we investigated whether ileal GLP-1R signaling is necessary for the glucose-regulatory effect of propionate (Fig. 4A). Infusion of exendin-9, a GLP-1R antagonist, alone into the ileal lumen had no effect on glucose metabolism, but co-infusion with propionate abolished the ability of the intraileal 50-min infusion of propionate to increase the glucose infusion rate (Fig. 4B), and lower GP (Fig. 4C, D). There were no changes in the rate of glucose uptake detected among the groups (Fig. 4E). In light of the fact that both ileal FFAR2 knockdown and GLP-1R inhibition negated the ability of propionate to lower GP and FFAR2 activation increases GLP-1 release, it is likely that GLP-1R signaling (hence the increase of GLP-1 levels) in the gut is a downstream mediator of the ileal propionate-FFAR2 axis in lowering GP. Future studies are warranted to address this working hypothesis.

As gut vagal afferent innervation is necessary for the metabolic effect of intestinal GLP-1 action (34), we next assessed the involvement of a neuronal network (Fig. 4A). Although the intraileal 50-min infusion of tetracaine, a local anesthetic, alone had no effect on glucose homeostasis, tetracaine negated the ability of the intraileal 50-min infusion of propionate to increase the glucose infusion rate (Fig. 4B) and decrease GP (Fig. 4, C and D), independent of changes in the rate of glucose uptake (Fig. 4E). These results indicate that ileal propionate activates ileal FFAR2 signaling to trigger a GLP-1R-dependent neuronal network to lower GP in vivo.

**Metabolism of LCFA in the ileum and a GLP-1 Receptor-dependent Neuronal Network Are Necessary for Ileal LCFA Sensing**—In addition to SCFAs, several studies suggest that the ileal mucosa comes in contact with a significant amount of ingested lipids (23, 29, 35). For example, 60 min following administration, ~12% of a radiolabeled IL load is present in the ileum (29). Therefore, given this, and the fact that intra-duodenal and -jejunal IL infusion triggers a neuronal-dependent negative feedback pathway to lower GP (14, 18), we first infused 5% IL into the ileal lumen of healthy rats for 50 min to activate ileal lipid sensing and evaluate the impact it has on glucose regulation. IL was infused at a dose one-fourth of that of previous duodenal studies (18), as IL is detected in a lower quantity in the ileum when compared with the duodenum following intragastric administration (29). Intraileal 50-min infusion of IL increased the glucose infusion rate (Fig. 5A) and lowered GP (Fig. 5, B and C), compared with saline vehicle alone infusion, with no changes detected in glucose uptake (Fig. 5D).

To further characterize ileal lipid sensing mechanisms, we next infused either the monounsaturated fatty acid, OA, or the...
polyunsaturated fatty acid, LA, solubilized in 1% bile acid (Fig. 6A), at a dose of 200 mM, which has previously been shown to stimulate gut peptide release in the ileum (23, 31). OA and LA were chosen based on the fact that IL is composed mainly of OA and LA, and that both LCFAs stimulate gut peptide release (36).

Independent intraileal 50-min infusion of either OA or LA increased the glucose infusion rate (Fig. 6B) and lowered GP (Fig. 6, C and D), compared with 1% bile acid vehicle alone infusion, with no change in glucose utilization (Fig. 6E) or plasma insulin levels (Table 1). This GP-lowering effect of OA or LA was preabsorptive, as plasma FFA levels were not elevated following OA, LA, or control in the intraileal 50-min infusions (Table 1), and intravenous infusions of OA or LA at the same dose and duration as intraileal LCFA infusion did not alter whole body glucose metabolism (Fig. 6, B–E). Our findings illustrate a novel ileal LCFA sensing pathway that lowers GP in vivo.

Esterification of duodenal and jejunal LCFAs into LCFA-CoA via acyl-C synthase is required to lower GP (14, 18). In the ileum, we here discovered that intraileal coinfusion of OA or LA with triacsin-C, an inhibitor of long-chain acyl-CoA synthase (Fig. 6A), abolished the ability of intraileal OA or LA 50 min infusion to increase the glucose infusion rate and lower GP, whereas triacsin-C + 1% bile acid vehicle alone into the ileum did not affect glucose metabolism (Fig. 6, B–E). These findings alternatively confirm that ileal LCFA preabsorptive sensing lowers GP and that metabolism of LCFAs in the ileum is a necessary step.

Last, we determined if an ileal GLP-1R-dependent neuronal network not only mediates the GP-lowering effect of ileal SCFA but also LCFA sensing (Fig. 6A). Intraileal infusion of exendin-9 + 1% bile acid vehicle alone had no effect on glucose metabolism (Fig. 6B) and decrease GP (Fig. 6, C and D). Furthermore, whereas tetracaine + 1% bile acid vehicle alone had no effect on glucose homeostasis (Fig. 6, B–E), co-infusion of tetracaine with OA or LA abolished the ability of the intraileal 50-min infusion of OA or LA to increase the glucose infusion rate (Fig. 6B) and decrease GP (Fig. 6, C and D). Thus, although the physiological relevance of ileal fatty acid sensing remains to be investigated, the current study suggests a mechanistic pathway for ileal fatty acids to decrease GP through a GLP-1R-dependent neuronal network in vivo.

**Discussion**

We here demonstrate, for the first time to our knowledge, that propionate sensing in the ileum triggers an ileal FFAR2 negative feedback pathway to lower GP through a GLP-1R-dependent neuronal network in vivo. This is in line with the fact that FFAR2−/− (but not FFAR3−/−) mice have reduced circulating GLP-1 levels that are associated with an impaired
FIGURE 4. GLP-1R signaling and a neuronal network are required for ileal propionate sensing. A, schematic representation of the working hypothesis. B–E, the glucose infusion rate (B), rate of GP (C), % suppression of GP (D), and glucose uptake (E) during the pancreatic clamp in rats with intraileal exendin-9 (Ex 9) and tetracaine (Tetra) administration with saline or in combination with propionate (n = 6 for all groups). Values are shown as mean ± S.E. **, p < 0.01; ***, p < 0.001, versus all other groups as determined by ANOVA with Tukey's post hoc test.

FIGURE 5. The effect of intraileal Intralipid infusion on glucose kinetics. A–D, the glucose infusion rate (A), rate of GP (B), % suppression of GP (C), and glucose uptake during the pancreatic clamp in rats with intraduodenal saline (Sal; n = 6) or Intralipid (IL; n = 6) infusions. Values are shown as mean ± S.E. **, p < 0.01; ***, p < 0.001, versus all other groups as determined by ANOVA with Tukey's post hoc test.
response to an oral glucose tolerance test (12), whereas the metabolic improvements of SCFA supplementation is maintained in FFAR3−/− mice (5). Our current discoveries not only directly integrate these associative findings for the first time, but also demonstrate site-specificity as the propionate-FFAR2-GLP-1 signaling axis in the ileum potently lowers GP in vivo. These findings highlight ileal FFAR2 as a potential mediator for the improved glucose homeostasis observed following SCFA supplementation either via direct intake of a SCFA-enriched meal or prebiotic treatment that increases SCFA production (5, 38, 39). Furthermore, given that gastric bypass surgery in rodents associates with a change in gut microbiota (i.e. stimulating SCFA production) and decrease in the acetate:propionate ratio (4), the currently described ileal propionate-FFAR2 signaling axis may also mediate the glucose-lowering effect of gastric bypass surgery, although these hypotheses clearly remain to be directly tested.

Of note, the ileal GLP-1 signaling pathway not only mediates propionate but also LCFA oleate and linoleate sensing in the ileum to lower GP, consistent with the fact that both LCFA and SCFAs stimulate GLP-1 release (13, 36). In addition, our discoveries illustrate that a neuronal network is necessary for ileal SCFA and LCFA sensing to lower GP independent of changes in plasma insulin levels. Thus, in addition to the well described incretin effect of GLP-1, our findings support an additional working hypothesis that suggest ileal fatty acids stimulate local GLP-1 release to act through an ileal GLP-1R-dependent paracrine neuronal axis to lower GP. This working hypothesis is supported by the fact that SCFAs, LCFA and GLP-1 trigger vagal firing (25, 40, 41), GLP-1Rs are expressed in the nodose ganglia that contains the cell bodies of vagal afferents (42), GLP-1 is rapidly degraded in circulation (43), and that the GLP-1R is in close proximity to L-cells (37). Although the direct site of vagal activation remains to be delineated (i.e. portal versus intestinal), our findings are in line with the notion that intestinal vagal afferent firing is necessary for nutrient-induced GLP-1 signaling to regulate glucose homeostasis (34). While the physiological relevance of ileal fatty acid sensing remains to be investigated, the current study suggests the presence of a mechanistic ileal fatty acid sensing pathway involved in glucose regulation.

FIGURE 6. Ileal LCFA metabolism and a GLP-1R dependent network are necessary for OA and LA sensing to lower glucose production. A, schematic representation of the working hypothesis. B–E, the glucose infusion rate (B), rate of GP (C), % suppression of GP (D), and glucose uptake (E) during the pancreatic clamp in rats with intraluminal or intravenous infusion of vehicle (1% bile acids), oleic acid (OA), or linoleic acid (LA), or intraluminal infusion of triacsin-C (Tri-C), exendin-9 (Ex 9), or tetracaine (Tetra) alone or coinfused with either OA or LA (n = 6 for all groups). Values are shown as mean ± S.E. **, p < 0.01; ***, p < 0.001, versus all other groups except LA or OA, as determined by ANOVA with Tukey’s post hoc test.
**Proposed pathway for ileal propionate sensing to lower glucose production.** In the ileum, microbiota-produced propionate activates FFAR2, localized in the enteroendocrine cells, and induces GLP-1 secretion. GLP-1 binds to GLP-1Rs located on the vagal afferent neurons, to ultimately reduce GP.

regulation. Interestingly, as more undigested LCFA could reach the ileum after bariatric surgery, future studies are warranted to address the relevance of ileal LCFA sensing and assess the relative contribution of jejunal (14) versus ileal nutrient sensing in the anti-diabetic effect of bariatric surgery.

In summary, we unveil that ileal short and long chain fatty acid sensing machinery triggers a GLP-1R-dependent negative feedback neuronal network to lower GP and regulate glucose homeostasis in rodents. Furthermore, we highlight FFAR2 in the ileum as a mechanistic target for lowering GP. Specifically, we postulate that microbiota-derived propionate activates FFAR2 in the ileal enteroendocrine cells to release GLP-1 and triggers local vagal GLP-1R to lower GP (Fig. 7). We propose changes in the fatty acid sensing mechanisms in the ileum regulate glucose homeostasis.

Author Contributions—M. Z-T. conducted and designed experiments, performed data analyses, and wrote the manuscript. F. A. D. edited the manuscript and conducted and assisted with experiments. B. A. R., P. V. B., C. D. C., and B. M. F. assisted with experiments. T. K. T. L. supervised the project, designed experiments, and edited the manuscript. T. K. T. L. is the guarantor of this work and, as such, had full access to all the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

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