Lactobacillus rhamnosus GG prevents epithelial barrier dysfunction induced by interferon-gamma and fecal supernatants from irritable bowel syndrome patients in human intestinal enteroids and colonoids

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ABSTRACT
Disruption of intestinal barrier homeostasis is an important pathogenic factor in conditions such as irritable bowel syndrome (IBS). Lactobacillus rhamnosus GG (LGG) improves IBS symptoms through unclear mechanisms. Previous studies utilizing colorectal adenocarcinoma cell lines showed that LGG metabolites prevented interferon gamma (IFN-gamma) induced barrier damage but the model employed limited these findings. We aimed to interrogate the protective effects of LGG on epithelial barrier function using human intestinal epithelial cultures (enteroids and colonoids) as a more physiologic model. To investigate how LGG affects epithelial barrier function, we measured FITC-Dextran (FD4) flux across the epithelium as well as tight junction zonula occludens 1 (ZO-1) and occludin (OCLN) expression. Colonoids were incubated with fecal supernatants from IBS patients (IBS-FSN) and healthy controls in the presence or absence of LGG to examine changes in gut permeability. Enteroids incubated with IFN-gamma demonstrated a downregulation of OCLN and ZO-1 expression by 67% and 50%, respectively (p<0.05). This was accompanied by increased paracellular permeability as shown by leakage of FD4. Pretreatment of enteroids with LGG prevented these changes and normalized OCLN and ZO-1 to control levels. These actions were independent of its action against apoptosis. However, these protective effects were not seen with LGG cell wall extracts, LGG DNA, or denatured (boiled) LGG. Intriguingly, IBS-FSN injected into colonoids increased paracellular permeability, which was prevented by LGG. LGG, likely due to secreted proteins, protects against epithelial barrier dysfunction. Bacterial-derived factors to modulate gut barrier function may be a treatment option in disorders such as IBS.

Introduction
The primary functions of the human gastrointestinal tract are to absorb nutrients and serve as a protective barrier against luminal contents including food antigens and microbes. The intestinal barrier is comprised of a single layer of epithelial cells connected by tight junction proteins that modulate paracellular permeability. Tight junctions are highly dynamic structures which open and close continuously in response to various stimuli. For example, cytokines, such as tumor necrosis factor-alpha (TNF-alpha) and interferon-gamma (IFN-gamma), play a crucial role in regulating tight junction protein expression and gut barrier function.

The gut microbiota comprises a diverse and complex community that closely interacts with the intestinal epithelium. There is a symbiotic host-microbe relationship with the microbiota providing essential functions, including antimicrobial protection as well as development and modulation of the gut immune system. The microbiota also play a critical role in regulating tight junction protein expression and interferon-gamma (IFN-gamma), play a crucial role in regulating tight junction protein expression and gut barrier function. Disruption of intestinal barrier integrity is an important factor in the pathogenesis of several highly prevalent and morbid diseases, including irritable bowel syndrome (IBS), inflammatory bowel disease (IBD). Furthermore, there is...
accumulating evidence that alterations in gut microbiota are a key pathogenic factor linked to gut barrier dysfunction, increased intestinal permeability, and inflammation in these disorders.\textsuperscript{19–22} Although it is unclear whether alterations in gut microbiota lead to gut barrier dysfunction or vice versa, considerable attention is focused on modifying these pathogenic factors as potential therapeutic options. However, the ability to modulate intestinal barrier function remains elusive.

Probiotics are defined as “live microorganisms that, when administered in adequate amounts, confer a health benefit on the host.”\textsuperscript{23} Probiotics may have beneficial actions on the host by excluding or inhibiting pathogens, enhancing epithelial barrier function, or by modulating host immune responses.\textsuperscript{24} Lactobacillus rhamnosus Gorbach-Goldin (LGG) is one of the most widely used and studied probiotics and has several biological properties that make it potentially useful as a probiotic. LGG is resistant to acid and bile, adheres well to the human intestinal epithelium, and produces factors with activity against many bacterial species.\textsuperscript{25} Studies suggest that LGG is important in promoting gut immune development, protection against inflammation-induced damage and stimulating gut barrier function.\textsuperscript{26–29} Clinically, LGG has been shown to improve IBS symptoms,\textsuperscript{30} delay onset of pouchitis after surgery for ulcerative colitis,\textsuperscript{31} and maintain remission in ulcerative colitis.\textsuperscript{32} Previous studies using epithelial colorectal adenocarcinoma cell lines showed that LGG prevented IFN-gamma induced epithelial barrier damage.\textsuperscript{26–28,33,34} However, cancer cell lines are unable to mimic normal physiology. For example, it has been reported that norovirus can only infect and replicate in human epithelial cells derived from organoids and not from cancer cell lines.\textsuperscript{35}

In this study, we employed human enteroids and colonoids, which are three-dimensional structures of human epithelium generated from isolated human intestinal crypts.\textsuperscript{36} We demonstrate that enteroids/colonoids are a physiologically relevant human model of gut barrier function and intestinal permeability that can be modulated by inflammatory cytokines. We implemented these systems to investigate whether pretreatment with LGG supernatant may prevent cytokine-evoked changes in tight junction protein expression and permeability in human enteroids. This may confirm or contradict findings using classical cell lines. To identify the component(s) of LGG responsible for stimulating tight junction protein expression, we investigated the effects of LGG extracted DNA, boiled LGG supernatant, LGG cell wall extract, and we compared the effect of LGG on tight junction protein expression and enteroid permeability to Lactobacillus crispatus. Lastly, we examined whether LGG may be useful for modulating gut barrier dysfunction seen in IBS by using human colonoids treated with fecal supernatants from diarrhea-predominant IBS (IBS-D) patients and healthy subjects in the presence or absence of LGG. We also investigated whether LGG regulates epithelial barrier function independent of its action against apoptosis.

Results

Human enteroid barrier function can be modulated by EGTA

We used human enteroids to evaluate paracellular barrier function under different conditions. FD4 was microinjected into the lumen of enteroids and images were obtained at different time points up to 20h. Under control conditions, the human enteroids retained 55\% of FD4 over 20h, while treatment of the enteroids with 2mM EGTA for 2h to disrupt tight junctions and impair permeability resulted in 7\% retention of the dye (Fig. 1). These results demonstrate that enteroids have an intact epithelial barrier, which can be modified by exogenous stimuli. We used this property of enteroids to measure epithelial barrier function and to monitor tight junction protein localization and mRNA expression over time in subsequent experiments.

IFN-gamma disrupts epithelial barrier function in a concentration-dependent manner

Recent studies showed that IFN-gamma released from the colonic mucosa of IBS patients was elevated compared to healthy controls.\textsuperscript{37} Hence we used IFN-gamma to disrupt epithelial barrier function. Previous in vitro studies used concentrations ranging from 10ng/ml to 100ng/ml of IFN-gamma to induce epithelial barrier dysfunction.\textsuperscript{34,38,39} The physiological implication of these concentrations...
is not known since we do not know the exact local concentrations of IFN-gamma in the mucosa. In the present study, we performed a dose-response study (60-500 ng/ml) to determine the concentration of IFN-gamma required to induce epithelial dysfunction in the enteroids. First, human enteroids were injected with FD4. Enteroids exposed to IFN-gamma showed a concentration- and time-dependent decrease in retention of FD4. Exposure to 500 ng/ml IFN-gamma for 20h resulted in a rapid loss of barrier integrity such that the fluorescence of the enteroids at 20h was 4% of the intensity at baseline. However, when enteroids were exposed to IFN-gamma at 60ng/ml, there was only 70% leakage of FD4 after 20h (Fig. 2A,2B).

We next evaluated gene expression of tight junction proteins in order to determine if IFN-gamma mediated disruption of barrier function is associated with altered tight junction expression. Incubation with increasing concentrations of IFN-gamma caused a progressive decrease in gene expression of zonula-occludens-1 (ZO-1) (p<0.05) (Fig. 2C). These results indicate that IFN-gamma causes a concentration-dependent decrease in tight junction gene expression and a corresponding increase in transepithelial permeability in human enteroids. IFN-gamma at a concentration of 200 ng/ml was chosen for subsequent experiments as this was the minimal concentration required for inducing reproducible epithelial barrier damage.

**LGG specifically protects against enteroid barrier dysfunction**

We next investigated whether modulation of barrier function induced by LGG was strain specific. To address this question, we compared its performance to Lactobacillus crispatus, another probiotic, known to prevent and treat recurrent bacterial vaginosis. Enteroids were precultured with either LGG-conditioned media (LGG-CM) or L. crispatus-CM overnight and then exposed to IFN-gamma 200 ng/ml for 24h. Enteroids in the presence of LGG-CM and IFN-gamma showed expression of occludin (OCLN) and ZO-1 similar to control levels. This was not seen in enteroids exposed to both L. crispatus-CM and IFN-gamma or IFN-gamma alone (Fig. 3A).
Consistent with qRT-PCR data, enteroids exposed to IFN-gamma caused a significant loss of barrier function whereas those exposed to both LGG-CM and IFN-gamma showed normalization of epithelial barrier function comparable to control levels. This rescue of barrier function was not seen in enteroids exposed to both L. crispatus-CM and IFN-gamma (Fig. 3B). This provided further evidence that bacterial components of LGG can inhibit cytokine-induced disruption of epithelial tight junction protein expression and intestinal mucosal barrier dysfunction. However, these beneficial actions were not seen in the presence of L. crispatus suggesting this protective property of LGG is species specific.

Proteins secreted by LGG prevents IFN-gamma induced epithelial barrier damage

We next investigated what components of LGG are responsible for preventing loss of mucosal barrier function. Enteroids were incubated for 24h with IFN-gamma with or without LGG-CM, LGG extracted DNA, boiled LGG-CM or LGG cell wall. Incubation of human enteroids with IFN-gamma caused 67% and 50% downregulation of OCLN and ZO-1 gene expression, respectively (p<0.05). Addition of LGG-CM prevented these changes and normalized OCLN and ZO-1 to control levels. However, addition of boiled LGG-CM, extracted LGG DNA or LGG cell wall abolished the protective effects of LGG-CM against IFN-gamma (Fig. 4A).

We then tested the permeability of the intestinal epithelium with or without LGG-CM, LGG extracted DNA, boiled LGG-CM or LGG cell wall. Under control conditions, the enteroids retained 70% of FD4 over 12h, while treatment of the enteroids with IFN-gamma impaired permeability resulting in 30% retention of dye after 12h. Administration of LGG-CM prevented leakage of dye evoked by IFN-gamma. In contrast, additions of extracted LGG DNA, cell wall or boiled supernatant were without effects (Fig. 4B).

We next performed immunofluorescence of the human enteroids. ZO-1 was normally present at the apical surface of the epithelium, whereas OCLN was located along the lateral/basal surface of the cell. Administration of IFN-gamma to human enteroids led to downregulation and disruption of ZO-1 and OCLN expression while pretreatment of enteroids with LGG-CM restored expression of ZO-1 and OCLN to control levels (Fig. 5). These observations indicate that protein metabolites secreted by LGG are responsible for preventing IFN-gamma induced epithelial barrier damage. This effect is mediated by normalizing the expression and localization of OCLN and ZO-1.
Figure 3. LGG specifically protects against human enteroid barrier dysfunction induced by IFN-gamma (A) Human enteroids were pretreated with LGG-CM or L. crispatus-CM overnight, and then they were exposed to IFN-gamma (200 ng/ml) for 24h. LGG-CM prevented IFN-gamma-induced downregulation of occludin and ZO-1 gene expression. However, this protective effect was not seen in enteroids incubated with L. Crispatus (*p<0.05, compared to control, CT). (B) Similarly, LGG-CM, but not L. Crispatus-CM, prevented leakage of the fluorescent dye FD4 induced by IFN-gamma. Under control conditions, the human enteroids retained 76% of FD4 at 12h. Treatment of the enteroids with IFN-gamma led to intestinal epithelial barrier dysfunction with only 35% of the dye retained at 12h. Administration of LGG, but not L. crispatus-CM, prevented leakage of dye evoked by IFN-gamma (*p<0.05, compared to control).
LGG regulates epithelial barrier function independent of its action against apoptosis

Previous studies using cancer cell lines showed LGG is capable of preventing apoptosis. We next investigated whether the protective action of LGG on epithelial barrier function is dependent on its action against apoptosis. Human enteroids were treated with IFN-gamma at 200ng/ml, or the “cytokine mixture” consisting of IFN-gamma (1000ng/ml) and TNF-a (1000 ng/ml) for 24h in the presence or absence of LGG-CM. Immunofluorescence study showed that cleaved caspase-3 (green fluorescence) was not expressed in enteroids exposed to IFN-gamma at 200ng/ml. Treatment with the cytokines...
TNF-a and INF-gamma at 1000ng/ml induced apoptosis which was not prevented by pretreatment of LGG-CM, indicating that the protective effect of LGG occurs independent of its action against apoptosis (Fig 6A, B).

The protective action of LGG on epithelial function is independent of MAPK/ERK pathway in human colonoids

Previous studies using cancer cell lines showed that LGG-derived soluble proteins protected against hydrogen peroxide-induced barrier dysfunction through a MAP kinase-dependent pathway. It is unknown whether a similar pathway is involved in the barrier protective actions of LGG in human colonoids. To examine this possibility human colonoids were treated with LGG-CM or epidermal growth factor (EGF) (100 ng/ml) for 90 min in the presence or absence of 1h pretreatment with (epidermal growth factor receptor) EGFR inhibitor, AG1478 (200nM). Cellular lysates were collected for Western blot analysis of total EGFR and extracellular signal-regulated kinase (ERK) levels and EGFR (Tyr-1068) and phosphorylated ERK. In human colonoids, LGG-CM treatment for 90min activated EGFR and its downstream effects. This was blocked by the EGFR inhibitor AG1478. In separate experiments, we examined whether the protective actions of LGG-CM on IFN-gamma induced barrier dysfunction was affected by inhibition of EGFR. As shown in Fig. 7, although AG1478 blocked the activation of EGFR and its downstream effects, it failed to affect the protective actions of LGG-CM on barrier dysfunction induced by IFN-gamma. This suggests that in humans, the actions of LGG on barrier function are mediated by mechanisms different from those observed in cell lines and animal models. Recently we showed that fecal supernatant of IBS-D patients impaired epithelial

Figure 5. Lactobacillus rhamnosus GG prevents downregulation of ZO-1 and occludin expression induced by IFN-gamma Human enteroids were pretreated with or without LGG-CM overnight, and then exposed to IFN-gamma (200 ng/ml). After treatment with IFN-gamma for 20 h, the human enteroids were fixed and stained. ZO-1 is present at the tight junction near the apical surface of the epithelium, whereas occludin stained in green is seen at the tight junction and along the lateral surface of the cell in control human enteroids. In contrast, human enteroids treated with IFN-gamma, apical ZO-1 at the tight junction is lost and occludin is no longer restricted to the lateral surface of the epithelial cell. Meanwhile, pretreatment of human enteroids with LGG-CM have immunofluorescence for ZO-1 and occludin similar to those of the control.
We next investigated whether LGG can prevent barrier dysfunction induced by fecal supernatant on IBS-D patients in the human colonoids.

**LGG protects against barrier dysfunction in human colonoids induced by fecal supernatants from IBS-D patients**

Fecal supernatants (FSN) from four IBS-D patients (IBS-FSN) as well as four healthy subjects were injected into human colonoids and retention of FD4 over time was measured. We observed little difference in the abilities to retain FD4 between colonoids from different healthy subjects. When exposed to FSN from healthy subjects (control) and IBS-D patients, the colonoids retained 80% of FD4 compared with 39% at 12 hours, respectively (p<0.05). Meanwhile, colonoids pretreated with LGG showed retention of FD4 similar to control levels, indicating a rescue of barrier function (Fig. 8).

**Discussion**

Although it has been demonstrated that LGG rescued barrier function and tight junction from cytokine-induced breakdown in mouse intestinal tissues and cancer cell lines, these models are unable to mimic normal physiology in humans. We demonstrated for the first time utilizing human enteroids and colonoids...
that secreted factors from LGG modulate epithelial barrier function and tight junction protein expression. We determined that barrier disruption induced by IFN-gamma was prevented by LGG-CM. We further showed that LGG normalizes tight junction expression and decreases mucosal permeability in the absence of apoptosis. In contrast to observations made in cancer cell lines, the action of LGG on barrier function is independent of activation of MAPK/ERK pathway. Finally, our results reveal that IBS-FSN disrupts intestinal barrier function in human colonoids which can be reversed by LGG. Our findings provide a molecular basis for therapeutic applications of LGG in gastrointestinal disorders, such as IBS-D.

Previous models of host-microbe interactions have utilized human epithelial colorectal adenocarcinoma cell lines, such as Caco-2 and HT29. A key difference in our model is the use of enteroids and colonoids, which are three dimensional structures derived from LGR5+ intestinal stem cells isolated from the small intestine or colon crypts rather than cancer-derived cell lines that are not physiologically relevant models of human intestinal function and structure. The human enteroids and colonoids have a single layer of epithelial cells with structural and planar cell polarity which assemble into polarized monolayers that separate central apical lumens from basal submucosa. This system is physiologically active which enables functional studies of tight junction expression and intestinal barrier function. We have confirmed the unique ability of enteroids and colonoids to study the complex host-microbe interactions.

We employed IFN-gamma to modulate tight junction protein expression and paracellular permeability. IFN-gamma is a pro-inflammatory cytokine that is elevated in the colonic mucosa of IBS patients and has been shown to directly decrease intestinal epithelial barrier function. We have demonstrated that IFN-gamma results in 40% reduction in gene expression of OCLN and ZO-1 in human enteroids which is consistent with previous studies.

Immunofluorescence was employed to confirm decreased expression as well as abnormal subcellular localization of OCLN and ZO-1. This was associated with increased paracellular permeability of enteroids by microinjection of FD4.

We further demonstrated that IFN-gamma induced disruption of gut epithelial permeability was effectively inhibited when enteroids were pretreated with LGG-CM. This was accompanied by
normalization of OCLN and ZO-1 gene expression. Our model suggests that LGG is unique in its ability to modulate epithelial barrier function as these protective effects were not seen when enteroids were pretreated with L. crispatus. L. crispatus is a probiotic capable of blocking uropathogens in vaginal epithelial cells and may prevent against recurrent urinary tract infections.45,46 In contrast to LGG, L. crispatus exacerbates murine colitis evoked by dextran sulfate sodium.47 Hence it was used as a negative control for our study.

Prior studies have demonstrated a cytoprotective effect of LGG via prevention against apoptosis. Yan et al. showed that LGG prevented cytokine induced apoptosis in intestinal epithelial cells through activation of Akt and inhibition of p38 activation.26 In contrast, our results indicate that LGG prevents cytokine-induced epithelial barrier damage by a mechanism independent from apoptosis.

LGG has been shown to be effective in treating gastrointestinal illnesses, such as acute gastroenteritis.48 However, the mechanisms by which LGG confer benefit are largely unknown. In addition, while probiotics are generally safe, there are potential risks in certain populations, such as immunocompromised or critically ill patients.49,50 Understanding the complex host-microbe interactions are still in its infancy, which is a major limitation for the prediction of efficacy, safety, and bioavailability of probiotics. One way to address these concerns is to understand how LGG interacts with the gut epithelium on a molecular level and then to isolate and purify the active factor(s) involved in these processes.

Although this study did not identify the specific factors responsible for LGG’s protective effects, we demonstrated that addition of extracted LGG DNA, LGG cell wall, and boiled LGG-CM did not alter intestinal epithelial permeability. This suggests that LGG exerts its effects on intestinal epithelial barrier function via secreted proteins. Prior studies have identified p40 and p75 as potential mediators of LGG-induced effects on epithelial
barrier function. These two novel proteins secreted by LGG may attenuate hydrogen peroxide-induced disruption of barrier function in Caco-2 cell monolayers. These protective effects likely occur via PI3K/Akt signaling pathway and MAPK-dependent signaling. Further studies have demonstrated that p40 ameliorates intestinal injury and colitis by stimulating ADAM17 activity and EGFR activation in colonic epithelial cells which stimulates mucin production through trans-activation of EGFR. Host-microbial interactions are complex and regulation of intestinal epithelial barrier function may be species and tissue specific. Actions observed in cancer cell lines may not be applicable to normal human epithelial lining. For example, we found that in contrast to cancer cell lines, the protective actions of LGG on tight junction proteins in human colonoids are not mediated by MAPK/ERK pathway. Furthermore, in human colonoids, LGG regulates epithelial barrier function independent of its action against apoptosis which is different from the observations made in HY-29 cell lines. Currently it is unclear whether p40 and p75 are responsible for normalizing expression of ZO-1 and OCLN. Future studies may determine whether p40 and p75 can be isolated from LGG-CM which may exert protective effects against inflammation-induced epithelial barrier dysfunction.

An intriguing finding in this study was that FSN taken from IBS-D patients, but not healthy controls, leads to impaired intestinal barrier function in human colonoids. However, pretreatment of colonoids with LGG prevented epithelial barrier dysfunction induced by IBS-FSN. De Palma et al. recently showed transfer of fecal samples from IBS-D patients to germ-free mice resulted in accelerated gastrointestinal transit and intestinal barrier dysfunction. Similar observations were made when fecal supernatant from IBS-D patients was administered into the colon of naïve rats. These findings indicate that soluble factors from IBS patients, possibly derived from gut microbiota, lead to pathophysiologic changes, including impaired gut barrier function. Furthermore, 39% of IBS-D patients have increased intestinal permeability as measured by the lactulose/mannitol ratio. These IBS patients demonstrate higher symptom scores and increased hypersensitivity to visceral nociceptive pain. Our findings indicate that LGG may directly improve mucosal barrier function by normalizing expression of junction proteins independent of immune modulation. This provides a rationale for using LGG to treat pain in IBS, which is a difficult symptom to treat. Future studies may investigate the potential of bacterial-derived factors to modulate intestinal barrier function in the treatment of conditions, such as IBS.

In conclusion, this study showed that LGG attenuates epithelial barrier dysfunction evoked by IFN-gamma in human enteroids. Furthermore, LGG normalizes tight junction protein expression which occurs in the absence of apoptosis. Our results demonstrate that protein metabolites secreted by LGG, but not bacterial DNA or cell wall, are responsible for preventing IFN-gamma induced epithelial barrier damage. Finally, our data indicate that IBS-FSN impaired intestinal barrier function in human colonoids, which was prevented by LGG. These findings support a potential application of bacterial components to prevent cytokine-mediated gastrointestinal injury and to treat epithelial barrier dysfunction in conditions, such as IBS. Identification of soluble factors mediating the beneficial effects of LGG may present an opportunity to understand their mechanism of action as well as to develop effective pharmacological strategies that may circumvent many of the problems posed by live bacterial therapies.

Methods

Human Specimens

Normal small intestinal and colonic tissue were obtained from patients undergoing surgical resection and colonoscopy, respectively, at the University of Michigan (UM). All human experiments were approved by the institutional review board at UM. Informed consent was obtained prior to acquisition of tissue.

LGG Conditioned Media Preparation

Lactobacillus rhamnosus GG (ATCC 53103) and Lactobacillus crispatus (ATCC 33820) (American Type Culture Collection (ATCC), Manassas, VA)
were incubated at 37 °C for 24h, then diluted in MRS broth according to ATCC guidelines.

For LGG-CM, LGG were inoculated in 50ml Dulbecco’s modified Eagle medium (DMEM) at 37°C overnight to reach log phase with the density determined as 0.6 at A600. The media was centrifuged twice, adjusted to pH 7.4, then filtered through a 0.2um filter to remove live bacteria.

**LGG cell wall isolation**

LGG was incubated in Lactobacillus MRS broth at 37 °C to reach log phase. Cultures were harvested by centrifugation and the cells were washed with PBS at room temperature twice. Cell suspensions were pipetted into 4% boiling SDS to lyse the cells for 3h. Boiled cells were ultracentrifuged (400,000 × g, 20 min, room temperature). The supernatant was removed and the pellets were resuspended in room temperature ultrapure water. Centrifugation was repeated and the samples were washed until SDS had been fully removed. The samples were resuspended in 10 mM Tris-HCl (pH 7.2) + 0.06% w/v NaCl. One mg/ml activated Pronase E (100 µg/ml final concentration) was added to each sample and incubated at 60 °C for 2h. Next, 200 µl of 6% SDS was added to stop the Pronase E digestion. Centrifugation and washing was repeated until the SDS had been fully removed. Samples were then re-suspended in 50 mM sodium phosphate buffer.

**LGG DNA Extraction**

LGG was incubated for 24 h at 37 °C in the exponential phase growth according to ATCC guidelines. After centrifugation (12000 rpm, 10 min) the bacterial pellets were used for total DNA extraction. These pellets were washed with NaCl-EDTA (30 mM NaCl, 2 mM EDTA, pH = 8.0) and resuspended in lysis buffer (Tris-HCl 20 mM, EDTA 2 mM, pH = 8.0), lysozyme (20 mg/mL) and triton X-100 (1% v/v). After incubation for 2h at 37°C, proteinase K (20 mg/mL) and RNase A (0.2 mg/mL) were added and incubated for 1h at 55°C. DNA was purified by repeat extraction with solvent of phenol-chloroform-isoamyl alcohol (25:24:1, v/v), precipitated with sodium acetate and ethanol, and dissolved in TE buffer. The purity of DNA was checked by a Nanodrop spectrophotometer.

**LGG boiled supernatant preparation**

LGG were inoculated in DMEM at 37°C to reach log phase with the density determined as 0.6 at A600. LGG-CM were denatured by boiling for 10 min.

**Human intestinal enteroid isolation and propagation**

Human intestinal tissue was washed with ice-cold Dulbecco’s Phosphate buffered saline without Ca2 + and Mg2+ (DPBS), and secured on a silicone-coated glass Petri dish filled with ice-cold DPBS. The mucosa was dissected from the underlying sub-epithelial tissue, washed 3–4 times with ice-cold chelation buffer to remove villi and debris, digested with freshly prepared 8mM EDTA chelation buffer for 30 min on a horizontal orbital shaker at 4°C. Epithelial crypts that were released from the tissue were collected by centrifugation of the media at 5 min at 800rpm, 4°C. Crypts were resuspended and then cultured in human enteroid complete medium (AdDMEM/F12 medium composed 50% LWRN conditioned medium, HEPES, Glutamax, penicillin/streptomycin, N2, B27, N-acetyl-L-cysteine, epidermal growth factor) mixed with Matrigel as previously described.43,58,59

The medium was replaced every other day.

**Enteroid treatment (IFN-gamma) and pretreatment (LGG-CM, L. crispatus-CM, cell wall, DNA, boiled LGG-CM)**

Three days after splitting, enteroids were cultured in human enteroid complete medium containing cell-free LGG-CM (5% vol/vol), LGG extracted DNA (10 µg/ml), boiled LGG-CM (5% vol/vol), LGG cell wall (4 mg/ml) or L. Crispatus-CM (5% vol/vol) overnight before treatment with IFN-gamma (200 ng/ml).
**Real-time quantitative polymerase chain reaction**

Cellular RNA was extracted from the enteroids by using Trizol Reagent (Invitrogen, Carlsbad, CA, USA) and Qiagen (Hilden, Germany) RNeasy mini columns, which was then reverse-transcribed into first-strand cDNA according to the manufacturer’s recommendations (iScript™ cDNA Synthesis Kit, Bio Rad, USA). The resultant cDNAs were used for RT-PCR, with primer sets targeting ZO-1 and OCLN. GAPDH served as an endogenous housekeeping reference gene.

**Immunofluorescent labeling**

Double or triple immunofluorescent staining was performed against ZO-1 and OCLN as well as Caspase-3, a key protease that is activated during the early stages of apoptosis. Enteroids were collected and fixed in 4% paraformaldehyde at 4°C for 30 min, then washed 3 times with PBS, and soaked in 30% sucrose for 24–48h at 4°C. Enteroids were embedded in optimal-cutting-temperature compound (4583; Sakura) for 20 min and frozen at -80°C. Frozen sections were cut at 8 µm for immunostaining, followed by microscopy. The cryostat sections were next rehydrated in PBS and blocked by 5% normal donkey serum (1:10; Chemicon International, Temecula, CA, USA) containing 0.3% Triton X-100 for 30 min at room temperature. The preparations were incubated overnight at room temperature with anti-ZO-1 antibody (1:500, ThermoFisher Scientific, USA), anti-occludin antibody (1:500, Invitrogen, USA) or caspase-3 (1:500, cell signaling). After incubation with the primary antibodies, the preparations were washed three times in PBS and incubated for 1h at room temperature with secondary antibodies Cy3 (1:200; Jackson ImmunoResearch Laboratories, West Grove, PA), Alexa 488-conjugated goat anti-rabbit IgG (1:200; Molecular Probes, Life Sciences Solutions)). Sections were then washed three times in PBS, mounted in buffered glycerol, and observed under fluorescence microscopy (Olympus BX-51, Tokyo, Japan). Images were captured at the same time of exposure, gain and gamma adjustment for the control and experimental groups.

**Human colonoids establishment**

Human colonic crypts were isolated from biopsy samples taken from healthy subjects undergoing colonoscopy at UM. The biopsy samples were washed 3–4 times with ice-cold PBS buffer containing penicillin-streptomycin (Pen/Strep, 1x), gentamicin (50 µg/ml), normocin (100 g/ml), and amphotericin (2.5 µg/ml) to control contamination, as well as thiazovivin (2.5 µM), a ROCK inhibitor. Crypts were isolated from biopsies by EDTA chelation containing 8 mM EDTA with DTT for 15 minutes, and in 8 mM EDTA for 15 minutes for further digestion. Crypts were collected in LWRN complete medium (AdDMEM/F12 medium composed of 50% LWRN conditioned medium, HEPES, Glutamax, penicillin/streptomycin, N2, B27, N-acetyl-L-cysteine, and epidermal growth factor). The crypt pellets were resuspended in basement membrane matrix and maintained in culture for 2 weeks.

**Western blot analysis**

To prepare total protein lysates, colonoids were harvested using BD Cell Recovery Solution in order to achieve Matrigel matrix depolymerization. The wells were washed with cold PBS and incubated for 60 minutes in cold BD Cell Recovery Solution. Once the colonoids were released, the cells were centrifuged at 800rpm for 5 minutes at 4°C. The pellet was homogenized in lysis buffer containing a mixture of proteinase and phosphatase inhibitors and then centrifuged at 15,000rpm for 15 min at 4°C. The protein concentration was determined using a BCA Protein Assay Kit (Pierce, Rockford, IL). Total protein was resolved on 4–12% precasted SDS-PAGE gels, then transferred to a polyvinylidene fluoride (PVDF) membrane (Millipore, Bedford, MA). The PVDF membrane was blocked with 5% non-fat milk in PBS containing 0.1% Tween 20 for 2 h at room temperature and then incubated overnight at 4°C with primary antibodies. The following antibodies were used in this study: GAPDH, anti-EGFR (phosphor Y1068) (1:2000, abcam, USA), total-EGFR, P-ERK, total-EGFR antibody (1:1000, cell signaling, USA). After washing with TBST, the blots were incubated for 2h at room temperature with HRP-conjugated
secondary antibody (1:5000; Amersham Biosciences, San Francisco, CA, USA), visualized by using Electro-Chemi-Luminescence (ECL) chemiluminescent detection system (Amersham Biosciences).

**Fecal supernatant preparation**

Four healthy subjects as well as four patients meeting Rome III criteria for IBS-D were recruited from outpatient clinics at UM. Subjects completed a 2 week screening period during which symptom severity was assessed based on 11-point numerical rating scale for abdominal pain, bloating and fecal urgency as well as stool consistency (Bristol stool form scale) and frequency. Six fecal samples were collected from each IBS-D patients and healthy subjects and stored at -80°C. Based on our recent studies, fecal samples were diluted (1 g fecal sample/5 ml PBS), homogenized on ice, and centrifuged (10,000 cpm, 10 minutes, 4°C). The supernatants were recovered, filtered on 0.22 μm filters to remove bacteria, and then stored at -80°C.

**Microinjection of Enteroids and Colonoids**

Each group of enteroids/colonoids was checked for integrity before injection. Thin-wall glass capillaries and tips were prepared as previously described. The capillaries were filled with 4kDa FITC-Dextran (FD4) and then loaded onto the microinjector (BRI XenoWorks analog microinjector; Sutter Instrument Company). Each enteroid/colonoid was injected with approximately 0.2 – 0.8ul of 1mMF4 based on the volume of enteroids, as described previously. Human enteroids and colonoids were imaged using a fluorescent stereomicroscope (SZX16; Olympus) at 1x magnification. Images were taken at the indicated time points postinjection. Disruption of barrier integrity was determined by loss of FD4 from the lumen of the enteroids/colonoids.

**Determining pixel intensity of FD4 in injected HIOs**

ImageJ software was used to determine the starting and final pixel intensity of the human enteroids and colonoids. Both bright-field and fluorescent images were taken from each well. Using the bright-field image, the perimeter of the enteroids or colonoids were outlined manually, and this region was used to determine the mean gray value of the enteroids or colonoids in the fluorescent image. These steps were repeated for each enteroid or colonoid for all treatments. The percent pixel intensity is defined as the mean gray value of an enteroid or colonoid at a given time point divided by the mean gray value of that same organoid at time 0 (T = 0) multiplied by 100.

**Statistical analysis**

All data were analyzed with SPSS 16.0 software (Chicago, IL). Differences between groups were compared by 2-tailed student’s t test or ANOVA for comparisons between 2 groups or more than 2 groups, respectively. Statistical significance was set at a P value of 0.05.

**Competing interests**

None declared.

**Contributors**

CO conceived and supervised the study. XU and CO designed and performed the experiments. XU, AL and CO wrote the manuscript. SH was involved in microinjection experiments. JG and JS were critical in human intestinal organoid culture and development. All authors discussed and interpreted the results.

**Disclosure of Potential Conflicts of Interest**

No potential conflicts of interest were disclosed.

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**Ethics approval**

All procedures were approved by the institutional review board (University of Michigan).

**Provenance and peer review**

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