

ORIGINAL ARTICLE

Food Allergy and Gastrointestinal Disease

Butyrate and propionate restore interleukin 13-compromised esophageal epithelial barrier function

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Funding information

NWO domain Applied and Engineered Sciences

Abstract

Background: Eosinophilic esophagitis (EoE) is a food allergen driven disease that is accompanied by interleukin (IL) 13 overexpression and esophageal barrier dysfunction allowing transepithelial food allergen permeation. Nutraceuticals, such as short-chain fatty acids (SCFAs) that restore barrier function and increase immune fitness may be a promising tool in the management of EoE. Here, we investigated the effects of the SCFAs acetate, propionate, and butyrate on an IL-13-compromised human esophageal epithelial barrier, including the mechanisms involved.

Methods: An air-liquid interface culture model of differentiated human EPC2-hTERT (EPC2) was used to study whether SCFAs could restore barrier function after IL-13-induced impairment. Esophageal epithelial barrier function was monitored by transepithelial electrical resistance (TEER) and FITC-dextran paracellular flux, and was further examined by qPCR and immunohistochemical analysis. G protein-coupled receptor (GPR) GPR41, GPR43, GPR109a, or histone deacetylase (HDAC) (ant)agonists were used to assess mechanisms of action of SCFAs.

Results: IL-13 stimulation decreased TEER and increased FITC flux, which was counteracted by butyrate and propionate, but not acetate treatment. Barrier proteins *FLG* and *DSG1* mRNA expression was upregulated following butyrate and propionate treatment, whereas expression of eosinophil chemoattractant *CCL26* and protease *CAPN14* was downregulated. Similarly, butyrate and propionate restored *FLG* and *DSG1* protein expression. Similar effects were observed with an HDAC antagonist but not with GPR agonists.

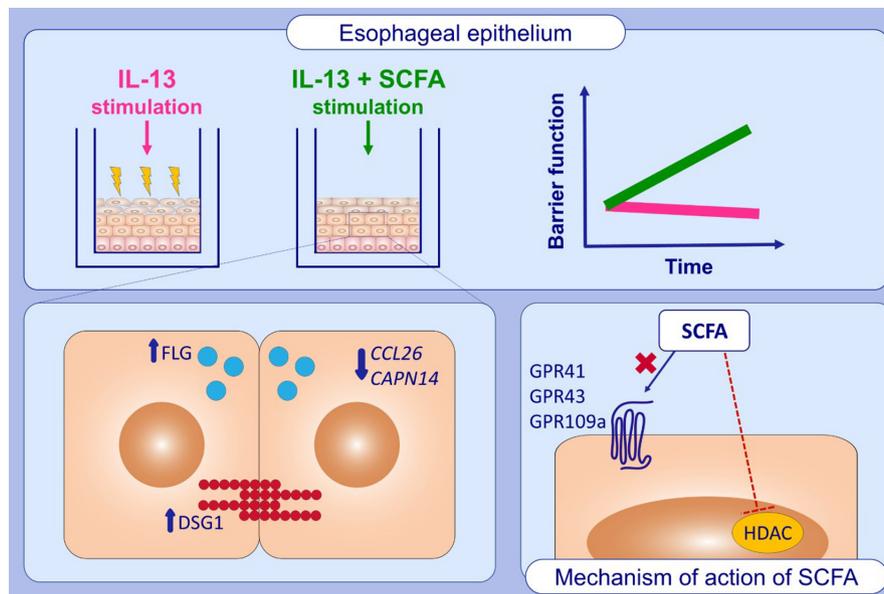
Conclusion: Nutraceuticals butyrate and propionate restore the barrier function of esophageal epithelial cells after an inflammatory insult and may be of therapeutic benefit in the management of EoE.

KEYWORDS

barrier function, dietary intervention, eosinophilic esophagitis, interleukin 13, short-chain fatty acids

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GRAPHICAL ABSTRACT

The SCFAs butyrate and propionate counteracted the compromising effects of IL-13 on barrier function of human esophageal epithelium cultured under ALI conditions. The increase in barrier function induced by these SCFAs was associated with restored expression of proinflammatory mediators and esophageal epithelial barrier proteins. An HDAC antagonist induced similar effects as butyrate and propionate, whereas GPR agonists did not.

Abbreviations: ALI, air-liquid interface; CAPN14, calpain-14; CCL26, (C-C motif) ligand 26; DSG1, desmoglein-1; FLG, filaggrin; GPR, G protein-coupled receptor; HDAC, histone deacetylase; IL-13, interleukin 13; SCFA, short-chain fatty acid.

1 | INTRODUCTION

The epithelial barrier of the esophagus forms the first line of chemical, physical, and immunologic defenses, and provides a protective wall against environmental factors including microbes and food allergens.¹ In eosinophilic esophagitis (EoE), a chronic food allergen-mediated disease of the esophagus, the esophageal barrier is frequently disrupted, leading to exposure to food allergens in the esophageal mucosa and the subsequent induction of a local type 2 immune response.^{2,3} Current treatment options for EoE consist of topical steroids and dietary restrictions^{4,5} but are sometimes unpopular with patients. Thus, there is a demand for novel treatment protocols that restore esophageal barrier function and mitigate esophageal inflammation to reestablish esophageal immune fitness.

Recent studies have demonstrated a link between the type 2 cytokine interleukin (IL) 13 in esophageal epithelial proliferation and esophageal barrier dysfunction.^{3,6,7} In fact, esophageal epithelial cells express each subunit of the IL-13 receptor including IL-4R α , IL-13R α 1, and IL-13R α 2.⁸ Transcriptomics studies have shown that IL-13 is overexpressed during active EoE, but its major cellular source or sources remain to be elucidated.⁸ Subsequently, IL-13 disrupts the esophageal barrier, mediated in part by the loss of the epithelial barrier proteins desmoglein-1 (DSG1) and filaggrin (FLG).^{3,9} In addition, IL-13 induces marked overexpression of eosinophil chemoattractant chemokine (C-C motif) ligand 26 (CCL26, encoding eotaxin-3) and protease calpain-14 (CAPN14).^{10,11} Notably, the EoE transcriptome can be partially reproduced in IL-13-treated immortalized esophageal epithelial cells cultured under air-liquid interface (ALI) conditions, indicating

that IL-13-induced gene expression in esophageal epithelial cells may make an important contribution to the EoE pathogenesis.⁶

Short-chain fatty acids (SCFAs) –in particular acetate, propionate, and butyrate– are produced by bacterial fermentation of dietary fiber in the gut, where they serve as an energy source for colonocytes, maintain intestinal homeostasis, and promote gut barrier function.¹²⁻¹⁴ SCFAs are agonists of G protein-coupled receptor (GPR) GPR41, GPR43, and GPR109a, inducing anti-inflammatory pathways upon binding.¹⁵⁻¹⁸ In addition, butyrate and propionate influence the activity of histone deacetylase (HDAC), a class of histone modification enzymes that regulates gene transcription and has the potential to influence biological processes.¹⁹⁻²² Although mainly produced in the gut, SCFAs have also been shown to have immunomodulatory effects in other barrier organs such as the lungs and skin.²³⁻²⁹

In this study, we use a model that resembles differentiated (ie, stratified squamous) human esophageal epithelium to investigate the potential barrier-restorative effects of the SCFAs acetate, propionate, and butyrate on an IL-13-compromised barrier. In addition, we aimed to determine the underlying mechanisms of the observed functional effects.

2 | MATERIALS AND METHODS

2.1 | EPC2-hTERT culture

The immortalized human esophageal epithelial cell line EPC2-hTERT (EPC2) was given by Dr Anil Rustgi (University of

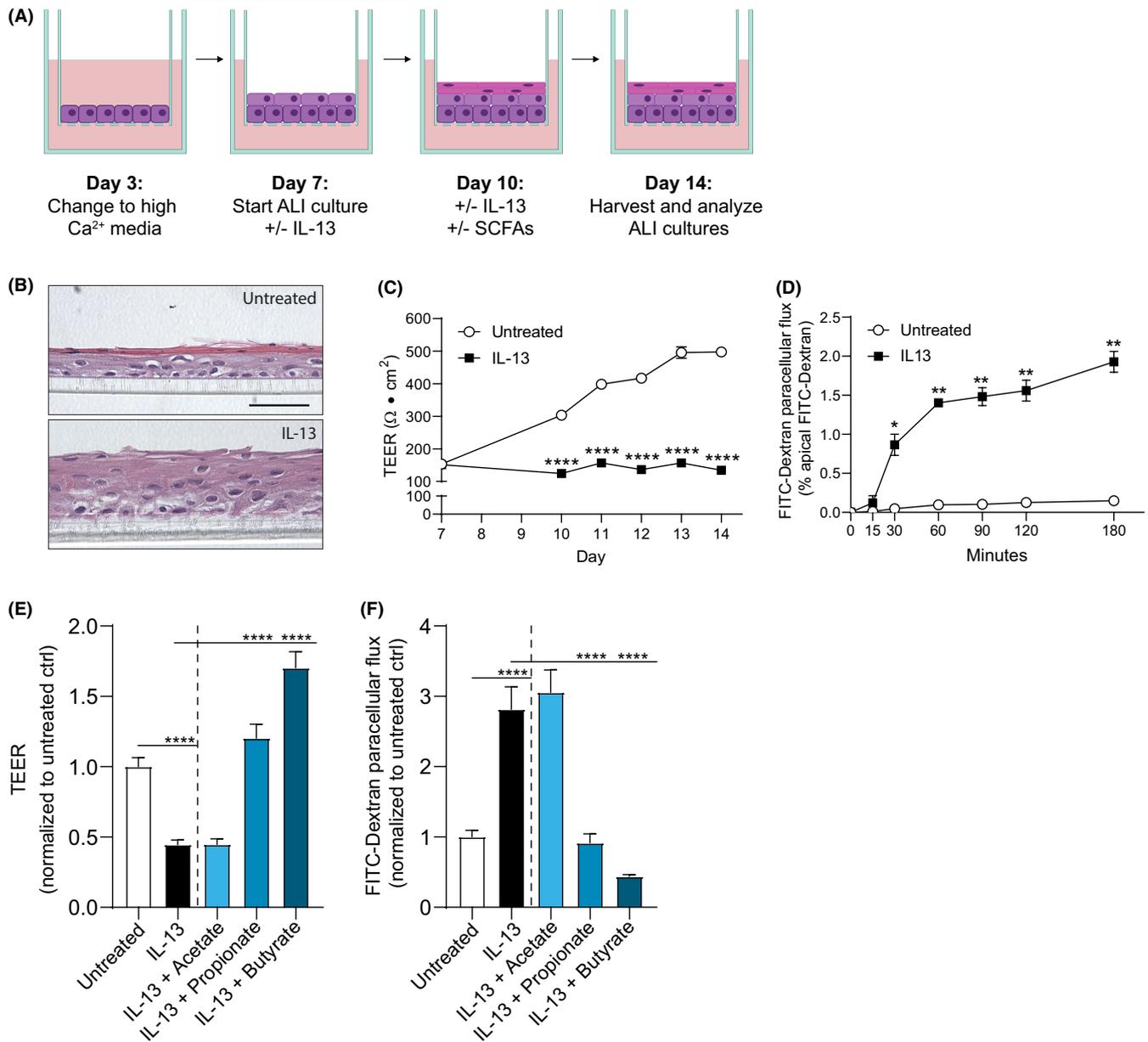


FIGURE 1 Butyrate and propionate restore IL-13-induced barrier dysfunction in EPC2 ALI cultures A, Schematic diagram of the ALI culture model. Culture day 1 to 7 allows initial differentiation, and culture day 7 to 14 (ALI) induces terminal differentiation and stratification of the EPC2. EPC2 are stimulated with IL-13 (100 ng/ml) from day 7 to 14. EPC2 are treated with SCFAs acetate (10 mM), propionate (10 mM), or butyrate (5 mM) from day 10 to 14. B, Hematoxylin and eosin staining of EPC2 differentiated at the ALI in the absence (untreated) or presence of IL-13 (100 ng/ml). Scale bar = 50 μm . C, TEER development of EPC2 in the absence (untreated) or presence of IL-13 (100 ng/ml) during differentiation under ALI conditions. D, Kinetic FITC flux analysis of EPC2 differentiated at the ALI in the absence (untreated) or presence of IL-13 (100 ng/ml). E and F, Day 14 TEER (E) and FITC flux (180 min.) (F) of IL-13-stimulated EPC2 ALI cultures treated with acetate (10 mM), propionate (10 mM) or butyrate (5 mM). Images and data in panels B-D are representative of twelve independent experiments ($n=6$ wells/group). Data in panels E and F are representative of two to eight independent experiments ($n=4$ wells/group). Data are presented as mean +SEM. Statistical significance was tested with one-way ANOVA followed by Dunnett's multiple comparisons test: * $p < .05$, ** $p < .01$, **** $p < .0001$

Pennsylvania, Philadelphia, PA, USA).³⁰⁻³² EPC2 were cultured in low-calcium (0.09 mM) keratinocyte serum-free medium (KSFM; Thermo Fisher Scientific, Waltham, MA, USA) supplemented with bovine pituitary extract (20 mg/ml), epidermal growth factor (1 ng/ml), penicillin (10,000 U/ml), and streptomycin (10,000 $\mu\text{g}/\text{ml}$).

2.2 | Air-liquid interface (ALI) culture system and SCFA treatment

The 3D ALI culture protocol was adapted from Kc et al.⁶ A schematic representation of the experimental timeline is shown in Figure 1A. Briefly, EPC2 were grown to confluence on semipermeable

membranes (0.4 μm ; Corning Incorporated, Corning, NY, USA) in low-calcium KSFM for 3 days. Initial differentiation of confluent monolayers was induced by switching to high-calcium (1.8 mM) KSFM from culture day 3 to 7. Terminal epithelial differentiation and stratification were induced by removing the media from the apical chamber and exposing the cells to the ALI from culture day 7 to 14. Cells were exposed to IL-13 (100 ng/ml; Prospec, Rehovot, Israel) in the basolateral chamber at the start of ALI culture.

Sodium acetate was purchased from BDH Laboratory Supplies (Poole, England, cat. no. 102364Q). Sodium propionate and sodium butyrate were purchased from Sigma-Aldrich (Saint-Louis, MO, USA, cat. no. P1880 (propionate) and 303410 (butyrate)). All SCFAs were used in preliminary work in a range of concentrations from 5 to 20 mM (acetate and propionate) and 2 to 10 mM (butyrate) (Figure S1). In final experiments, acetate (10 mM), propionate (10 mM), or butyrate (5 mM) was added to the basolateral chamber of IL-13-stimulated EPC2 ALI cultures from day 10 to 14. EPC2 ALI cultures were also treated with SCFAs in the absence of IL-13 (Figure S2). Media plus IL-13 and SCFAs were refreshed every other day. ALI cultures were then collected for total RNA isolation and immunohistochemistry.

2.3 | Assessment of mechanisms of action of SCFAs

The following (ant)agonists were used to investigate the involvement of GPR41, GPR43, GPR109a, and HDAC in the barrier-restorative effects of SCFAs: GPR41 agonist AR420626 (1 μM), GPR43 agonist 4-CMTB (10 μM), GPR109A agonist niacin (10 mM), and HDAC antagonist trichostatin A (TSA, 2 μM). All (ant)agonists were dissolved in DMSO or 1 M NaOH according to the manufacturer's instructions and were purchased from Sigma-Aldrich (cat. no. SML1339 (AR420626), SML0302 (4-CMTB), N4126 (niacin), and T8552 (TSA)). All (ant)agonists were used in preliminary ALI experiments in a range of concentrations from 1 to 100 μM (AR420626), 0.1 to 50 μM (4-CMTB), 1 to 20 mM (niacin), and 0.1 to 10 μM (TSA) (Figures S4 and S5). In final experiments, (ant)agonists were added to the basolateral chamber of the IL-13-stimulated EPC2 ALI cultures from day 10 to 14. Media plus IL-13 and (ant)agonist were refreshed every other day. ALI cultures were then collected for total RNA isolation and immunohistochemistry.

2.4 | Transepithelial electrical resistance (TEER), paracellular flux assays and LDH toxicity test

TEER was measured during ALI culture using a Millicell ERS-2 Volt-ohm meter (Merck Millipore, Burlington, MA, USA). High-calcium KSFM was added to the apical chamber one hour before TEER measurement. Paracellular flux assays were performed one hour after TEER measurement on day 14. 4-kDa fluorescein isothiocyanate (FITC)-dextran (0.1 mg/ μL ; Sigma-Aldrich) was added to the apical chamber, and fluorescein levels in the basolateral chamber were detected after 15, 30, 60, 90, 120, and 180 minutes using a GloMax

Discover Microplate Reader (Promega, Madison, WI, USA) at Ex/Em = 492/518. Cytotoxicity was measured in 50 μL supernatant collected at day 14 using the Cytotoxicity Detection Kit (LDH) (Roche, Basel, Switzerland) (Figure S7) per manufacturer's instructions.

2.5 | Quantitative real-time PCR

Total RNA was treated with DNase I (Qiagen, Hilden, Germany) and isolated from EPC2 ALI cultures using the RNeasy Mini Kit (QIAGEN) according to the manufacturer's instructions. cDNA was generated from 500 ng RNA using the iScript™ cDNA Synthesis Kit (BioRad, Hercules, CA, USA). qPCR was performed using SYBR Green (BioRad). All primers used for amplification were purchased from BioRad (Unique Assay ID: qHsaCID0017001 (CAPN14), qHsaCED0041923 (CCL26), qHsaCED0044569 (DSG1), and qHsaCED0036604 (FLG)). Results were normalized to ribosomal protein S13 (RPS13; Unique Assay ID: qHsaCID0038672) expression for each sample. mRNA expression levels were calculated using the following formula: fold change = $2^{-\Delta\Delta\text{Ct}}$ and were normalized to the untreated control.

2.6 | Histology and Immunofluorescence

Formalin-fixed, paraffin-embedded EPC2 ALI cultures were cut into 5 μm sections and deparaffinized using xylene followed by graded ethanol washes. For histology, sections were stained in hematoxylin, rinsed in tap water and then stained in eosin, followed by dehydration in graded ethanol washes and xylene before mounting with Pertex (Histolab, Askim, Sweden) and xylene (1:1). For immunofluorescence, heat-induced antigen retrieval in sodium citrate buffer (10 mM citric acid, pH 6.0) was used on deparaffinized sections, and endogenous peroxidase activity was quenched using 3% H_2O_2 in methanol. After rinsing in 0.2% Tween in PBS, sections were blocked in 3% BSA in PBS containing 5% normal goat serum (Dako, Jena, Germany) for 90 minutes, followed by overnight incubation at 4°C with rabbit anti-DSG1 (1 $\mu\text{g}/\text{ml}$; Abcam, Cambridge, UK, cat. no. ab209490) or rabbit anti-FLG (1 $\mu\text{g}/\text{ml}$; Abcam, cat. no. ab234406). Sections were rinsed and incubated for 1 hour with goat anti-rabbit AF594 (10 $\mu\text{g}/\text{ml}$; Invitrogen, Carlsbad, CA, USA, cat. no. A11072). Sections were mounted with ProLong™ Gold antifade reagent with DAPI (Invitrogen) for nuclei staining. Immunofluorescent images were acquired using the Keyence Fluorescence Microscope BZ-9000, and immunofluorescence intensity was quantified using ImageJ software.

2.7 | Nuclear extract preparation and HDAC activity

EPC2 were grown in 12 well culture plates (Costar) in low-calcium KSFM until confluent, followed by stimulation with acetate (10 mM), propionate (10 mM), butyrate (5 mM), or TSA (2 μM) in high-calcium KSFM. Cytoplasmic and nuclear extracts were isolated 48 hours after

stimulation. Briefly, EPC2 were trypsinized, collected by centrifugation (1000 rpm, 4 min, 4°C) and washed twice in ice-chilled PBS. EPC2 were resuspended in 100 μ L ice-chilled buffer 1 (Table S1) and incubated on a rotator for 10 minutes at 4°C. After vortexing, lysates were spun down (12,000 rpm; 1 min; 4°C) and the cytoplasmic protein fractions were collected and stored at -80°C. Nuclear pellets were washed twice with ice-chilled PBS, disrupted with 40 μ L Buffer 2 (Table S1) and incubated on ice for 30 minutes with regular vortexing followed by sonication for 3 x 10 seconds. The suspension was spun down (12,000 rpm; 15 min; 4°C) and the nuclear fractions were collected and stored at -80°C. Total protein content was quantified using the Pierce™ BCA Protein Assay Kit (Thermo Fisher Scientific). HDAC activity was measured in 2 ng nuclear extract using the colorimetric epigenase HDAC Activity/Inhibition Direct Assay Kit (EpiGentek, Farmingdale, NY, USA) according to the manufacturer's instructions.

3 | RESULTS

3.1 | Short-chain fatty acids butyrate and propionate restored esophageal barrier resistance and permeability in IL-13-stimulated EPC2 ALI cultures

We used an ALI culture model that resembles human-differentiated esophageal epithelium to investigate the potential barrier-restorative effects of SCFAs as depicted schematically in Figure 1A. Following 7 days of differentiation at the ALI, EPC2 formed a stratified squamous epithelial layer indicating the development of differentiated esophageal epithelium (Figure 1B). Prolonged IL-13 exposure resulted in marked morphologic changes including decreased epithelial differentiation and expansion of the epithelial layer (Figure 1B). Furthermore, IL-13 induced a significant decrease in TEER from day 10 and onwards (Figure 1C), and a significant increase in FITC-dextran paracellular flux (FITC flux) on day 14 (Figure 1D). Together, these results indicate that IL-13 induces barrier dysfunction in EPC2 ALI cultures as reported previously.^{3,6}

To study the barrier-restorative effects of SCFAs on IL-13-stimulated EPC2 ALI cultures, acetate, propionate, or butyrate were added to the basolateral chamber from day 10 to 14. At day 14, IL-13 stimulation showed a 2.2-fold decrease in TEER compared with untreated cultures. Propionate and butyrate counteracted the effect of IL-13 on TEER as shown by a 2.7-fold increase in propionate-treated ALI cultures and 3.8-fold increase in butyrate-treated cultures compared with IL-13-stimulated EPC2 ALI cultures (Figure 1E). FITC flux assays confirm these findings. IL-13-stimulated EPC2 ALI cultures had a significantly increased FITC flux at day 14, which was counteracted by propionate and butyrate treatment (Figure 1F). In addition, SCFAs -in particular butyrate- restored IL-13-induced barrier dysfunction measured by TEER and FITC flux in a culture model of apical SCFA treatment, supporting our data on basolateral SCFA treatment (Figure S3). Together, these data show that butyrate and propionate, but not acetate, restore esophageal barrier resistance and permeability after IL-13-induced impairment.

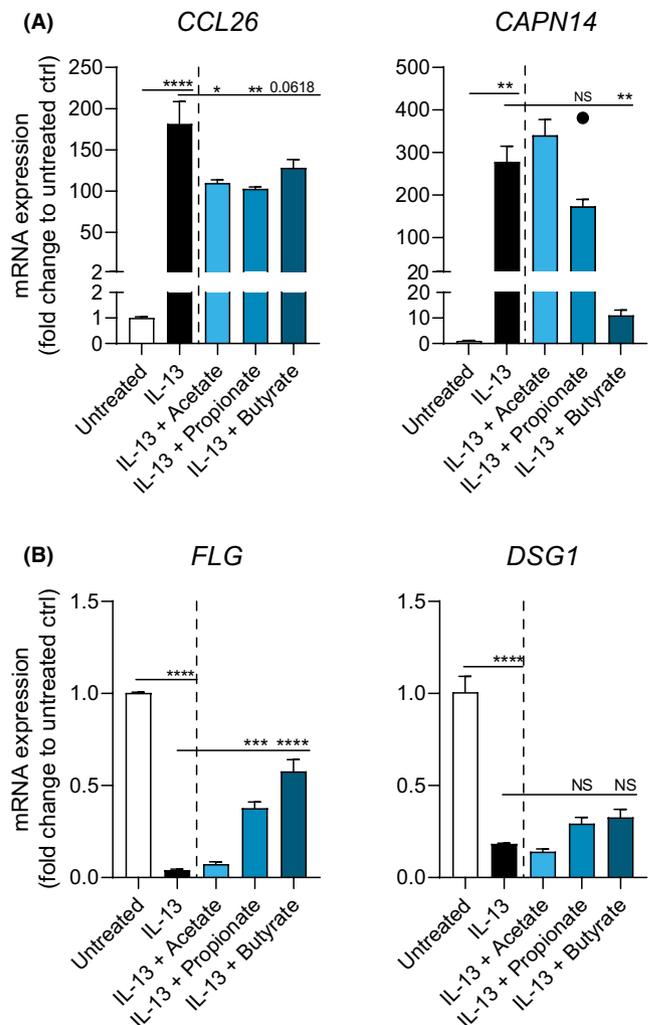


FIGURE 2 Butyrate and propionate restore mRNA expression of EPC2 ALI cultures. A, mRNA expression of proinflammatory factor *CCL26* and protease *CAPN14* in IL-13-stimulated EPC2 ALI cultures treated with acetate (10 mM), propionate (10 mM), or butyrate (5 mM). B, mRNA expression of esophageal barrier proteins *FLG* and *DSG1* in IL-13-stimulated EPC2 ALI cultures treated with acetate (10 mM), propionate (10 mM), or butyrate (5 mM). Data are representative of two to six independent experiments (n=3 wells/group). Outlier is shown as a separate data point. Data are presented as mean +SEM. Statistical significance was tested with one-way ANOVA followed by Dunnett's multiple comparisons test: * $p < .05$; ** $p < .01$; *** $p < .001$; **** $p < .0001$; NS, not significant

3.2 | Butyrate and propionate restored mRNA expression of key EoE genes

qPCR analysis was used to assess whether SCFAs changed mRNA expression of proinflammatory factor *CCL26*, protease *CAPN14*, and barrier proteins *DSG1* and *FLG*. IL-13 treatment significantly increased *CCL26* and *CAPN14* mRNA expression and significantly decreased *DSG1* and *FLG* mRNA expression by day 14. This was counteracted by propionate and butyrate as they decreased the expression of *CCL26* and *CAPN14*, while increasing the expression of *FLG* and *DSG1* compared with IL-13-stimulated EPC2 ALI cultures

(Figure 2A, B). These results correspond with the observed improved barrier function after butyrate and propionate treatment and further indicate that treatment with these SCFA have an anti-inflammatory action.

3.3 | Butyrate and propionate restored DSG1 and FLG protein expression

To test the effects of SCFA treatment on esophageal barrier protein expression, we examined day 14 DSG1 and FLG expression by

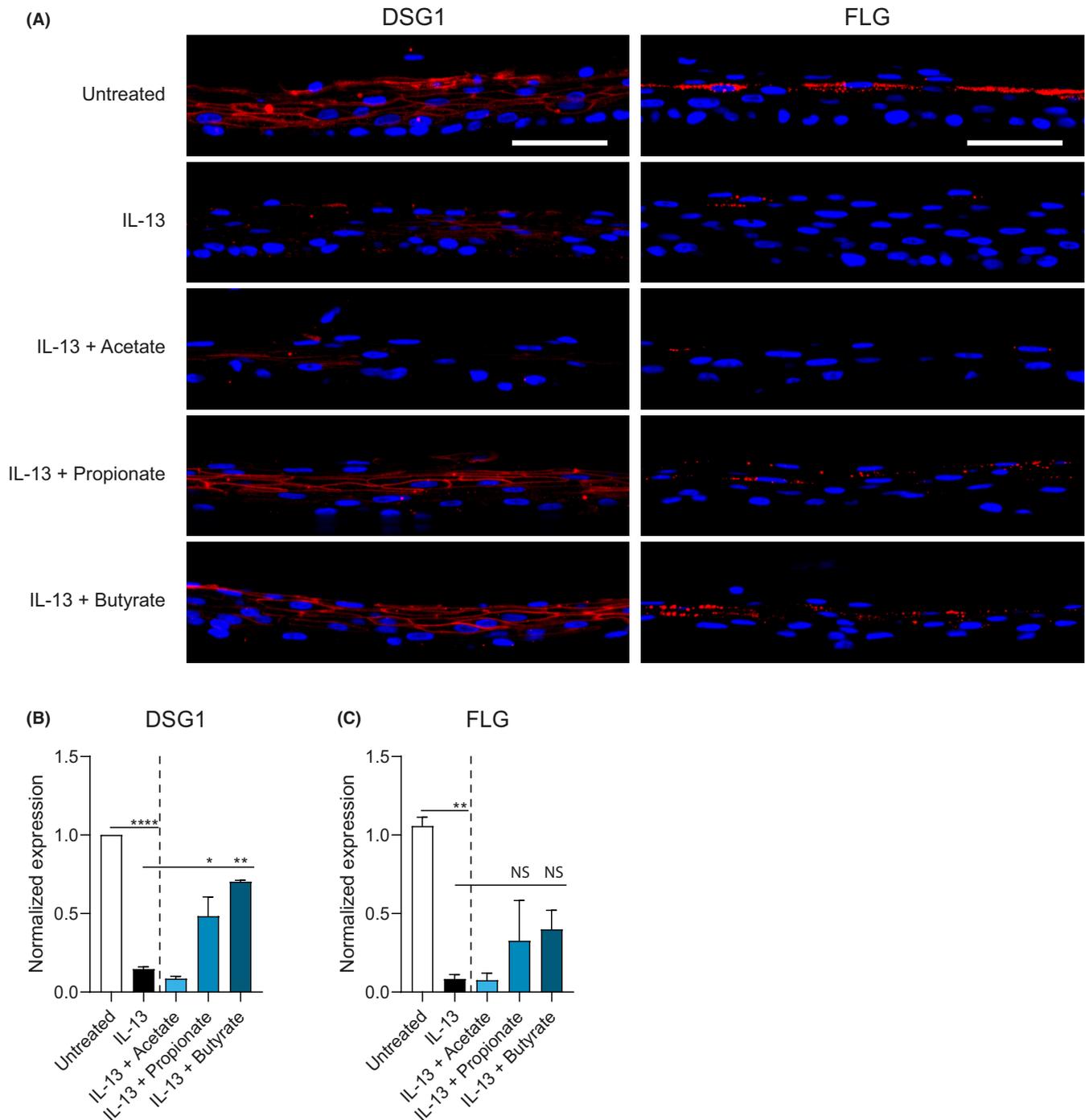


FIGURE 3 Butyrate and propionate upregulate DSG1 and FLG protein expression in ALI cultures of EPC2 treated with IL-13. A, Immunofluorescent staining for barrier proteins DSG1 (left) and FLG (right) in red with a blue DAPI nuclear counterstain in IL-13-stimulated EPC2 ALI cultures treated with acetate (10 mM), propionate (10 mM), or butyrate (5 mM). Scale bar = 50 μ m. B, Quantification of DSG1 and FLG expression in IL-13-stimulated EPC2 ALI cultures treated with SCFA. Images in panel A are representative of three independent experiments performed in duplicate or triplicate and are taken at 40x magnification. Data in panel B are pooled from three independent experiments performed in duplicate or triplicate and are presented as mean \pm SEM. Statistical significance was tested with one-way ANOVA followed by Dunnett's multiple comparisons test: * p <.05, ** p <.01; NS, not significant

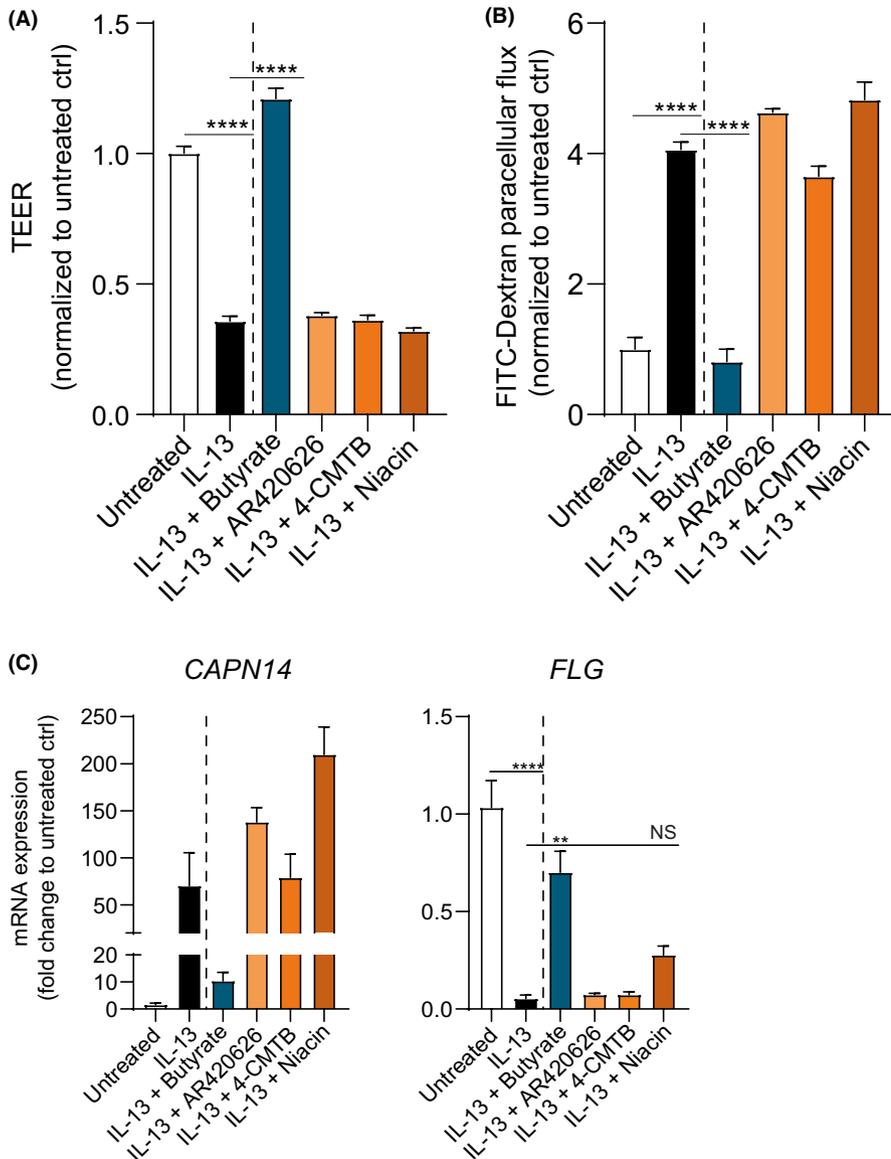


FIGURE 4 The effects of SCFAs are independent of GPR41, GPR43, and GPR109a stimulation. Day 14 TEER (A) and FITC flux (180 min.) (B) of IL-13-stimulated EPC2 ALI cultures treated with GPR agonists AR420626 (1 μ M, GPR41), 4-CMTB (10 μ M, GPR43), or niacin (10 mM, GPR109a). C, CAPN14 and FLG mRNA expression in IL-13-stimulated EPC2 ALI cultures treated with GPR agonists. Data are representative of two independent experiments (n=3 wells/group) and are presented as mean +SEM. Statistical significance was tested with one-way ANOVA followed by Dunnett's multiple comparisons test: ** $p < 0.01$; *** $p < 0.001$; **** $p < 0.0001$; NS, not significant

immunofluorescent staining. DSG1 and FLG expression was decreased in IL-13-stimulated EPC2 ALI cultures compared with untreated ALI cultures. Consistent with the mRNA expression data, butyrate, and to a lesser extent propionate, restored the expression of DSG1 and FLG in IL-13-stimulated EPC2 ALI cultures (Figure 3A). Quantification of fluorescence intensity confirms the upregulation of DSG1 and FLG in IL-13-stimulated EPC2 ALI cultures after butyrate and propionate treatment (Figure 3B, C).

3.4 | Barrier-restorative effects of butyrate and propionate are independent of the free fatty acid receptors GPR41, GPR43, and GPR109a

Next, we investigated whether the barrier-restorative effects of butyrate and propionate depend on signaling through the free fatty acid receptors GPR41, GPR43, and GPR109a. All three GPRs were

found expressed in EPC2 on mRNA and protein level (data not shown). Direct stimulation of GPRs with specific agonists did not affect neither TEER (Figure 4A) nor FITC flux (Figure 4B). In line with these observations, expression of genes associated with EoE and altered by IL-13 stimulation of EPC2 grown under ALI conditions was unaffected by GPR stimulation (Figure 4C).

To confirm the ability of these agonist to stimulate GPRs and reduce inflammation, human umbilical vein endothelial cells (HUVECs) were stimulated with lipopolysaccharide (LPS) following treatment with GPR agonists. Stimulation of GPRs with specific agonists decreased LPS-induced IL-6 and IL-8 release in a concentration-dependent manner (Figure S6), confirming the ability of the GPR agonists used in this study to stimulate GPR signaling. Together, these data indicate that the barrier-restorative effects of butyrate and propionate measured by TEER, FITC flux, and mRNA and protein expression are most likely not mediated via stimulation of GPR41, GPR43, or GPR109a in EPC2.

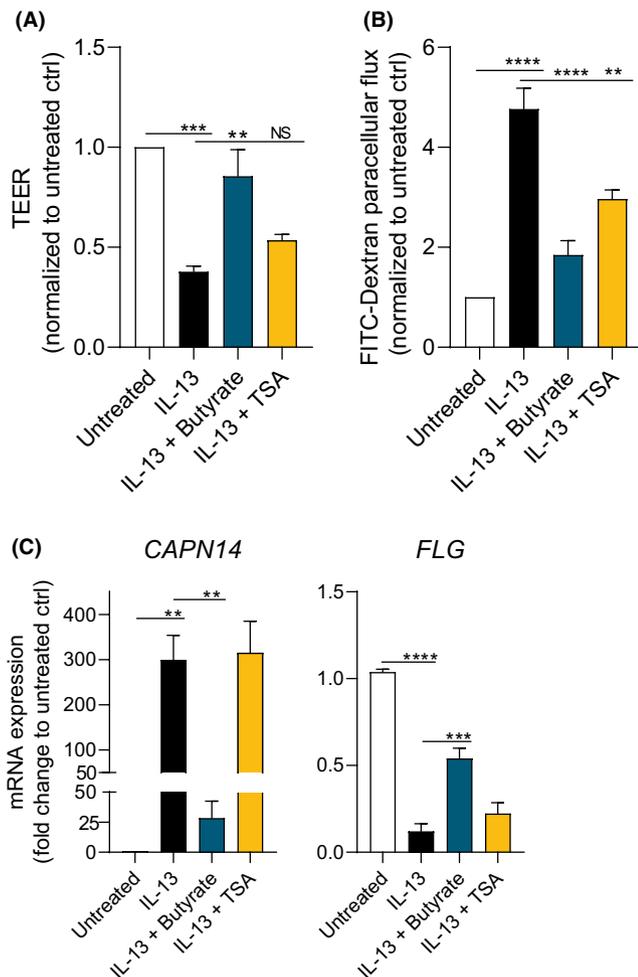


FIGURE 5 The HDAC inhibitor TSA partially mimics effects in EPC2 ALI cultures. Day 14 TEER (A) and FITC flux (180 min.) (B) of IL-13-stimulated EPC2 ALI cultures treated with TSA (2 μ M). C, *CAPN14* and *FLG* mRNA expression in IL-13-stimulated EPC2 ALI cultures treated with TSA (2 μ M). Data are pooled from four independent experiments performed in duplicate, triplicate, or quadruplicate and are presented as mean \pm SEM. Statistical significance was tested with one-way ANOVA followed by Dunnett's multiple comparisons test: ** $p < .01$; *** $p < .001$; **** $p < .0001$; NS, not significant

3.5 | HDACs may be involved in barrier-restorative effects of butyrate and propionate

It has been demonstrated that SCFAs are also effective inhibitors of HDAC activity.^{19,21,22} Since the effects of butyrate and propionate are independent of GPR signaling, we investigated whether the barrier-restorative effects of SCFAs may be related to inhibition of HDAC. To study the functional effects of HDAC inhibition, TSA, a potent and specific inhibitor of HDAC activity was added to EPC2 ALI cultures. Despite the minimal effect on TEER (Figure 5A), TSA significantly decreased FITC flux in IL-13-stimulated EPC2 ALI cultures, although with a smaller impact than butyrate (Figure 5B). *CAPN14* and *FLG* mRNA expression was not affected by TSA (Figure 5C). Furthermore, butyrate and propionate treatment led to attenuated HDAC activity in EPC2 (Figure 6). These data suggest that inhibition

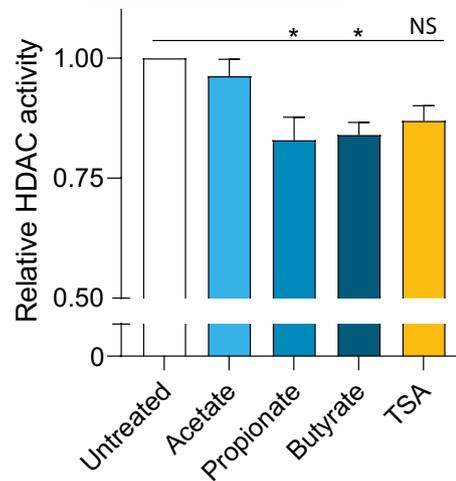


FIGURE 6 Butyrate and propionate decrease HDAC activity in EPC2. HDAC activity was measured in 2 ng nuclear proteins after treating confluent EPC2 for 48 hours with acetate (10 mM), propionate (10 mM), butyrate (5 mM), or TSA (2 μ M) and was normalized to the untreated control. Data are pooled from three independent experiments performed in triplicate, and are presented as mean \pm SEM. Statistical significance was tested with one-way ANOVA followed by Dunnett's multiple comparisons test: * $p < .05$; **** $p < .0001$

of HDAC activity can partly mimic the restorative effects on epithelial barrier function as observed by butyrate and propionate.

4 | DISCUSSION

The data presented in this study demonstrate that the SCFAs butyrate and propionate, but not acetate, restore esophageal epithelial barrier function after IL-13-induced impairment using an ALI culture model resembling differentiated human esophageal epithelium. First, we demonstrate that butyrate and propionate restore epithelial barrier resistance and permeability, as assessed by TEER and FITC flux. Second, we show that butyrate and propionate restore mRNA expression of genes associated with inflammation in EoE, such as *CCL26*, and barrier function, such as *CAPN14*, *DSG1*, and *FLG*. Third, we show that butyrate and propionate increase *DSG1* and *FLG* protein expression. Fourth, our studies suggest that the barrier-restorative effects of butyrate and propionate are independent of GPR signaling, but may, in part, be dependent on inhibition of nuclear HDAC activity.

Although acetate is the most abundant SCFA in the gut and periphery, butyrate is the most potent immunomodulatory SCFA.³³ Indeed, we observed that butyrate has the highest potency to enhance esophageal barrier function after IL-13-induced impairment. Also propionate, but not acetate, significantly augmented barrier function despite with a lower activity than butyrate. Our data add to the growing body of literature linking SCFAs to immunomodulation and epithelial barrier function. Nonetheless, Wen et al. have reported a potential proinflammatory effect of SCFAs in Th2 cell-associated responses,³⁴ indicating that

immunomodulatory effects of SCFAs are cell type-dependent. Our findings are consistent with effects of SCFAs on cytokine-compromised monolayers of Caco-2 and T84 human colorectal carcinoma cells and 16HBE human bronchial epithelial cells, where butyrate enhanced barrier function and tight junction protein expression at millimolar level.³⁵⁻³⁸ We used relatively high SCFA concentrations compared with these studies, which could be attributed to characteristics of the stratified esophageal epithelial layer that may contribute to SCFA sensitivity.

Here, we focused on the response of CAPN14 protease and esophageal barrier proteins DSG1 and FLG expression to SCFA treatment because of their suggested role in esophageal barrier function.^{3,9,11,39} Expression of the epithelium-derived proinflammatory factor CCL26 was studied because of its strong correlation with disease severity.¹⁰ The increase in TEER and decrease in FITC flux induced by butyrate and propionate were associated with a decrease in mRNA expression of *CCL26* and *CAPN14*, and an increase in mRNA and protein expression of DSG1 and FLG. CAPN14 activity is specific for esophageal tissue and its overexpression results in loss of epithelial barrier function.^{11,40-42} Furthermore, previous studies have shown that DSG1 and FLG are downregulated in inflamed esophageal mucosa of EoE patients,^{3,6,9} but are restored after successful therapeutic treatment and are associated with improved mucosal integrity.^{43,44} Whereas, DSG1 is specifically linked to EoE pathology,³ IL-13-mediated downregulation of FLG has also been described in atopic dermatitis.^{45,46} The role of other epithelial barrier proteins including claudins, occludin, involucrin, E-cadherin, and keratins in maintaining esophageal epithelial integrity is less evident.^{6,47} Interestingly, rather than changes in tight junction proteins, DSG1 and FLG dysregulation contributes to esophageal barrier dysfunction.⁴⁴ Current findings indicate that SCFAs can restore dysregulated expression of DSG1 and FLG leading to restoration of esophageal barrier function. In addition to IL-13, transforming growth factor (TGF) β 1 and IL-9 have also been found to diminish esophageal barrier function of esophageal epithelial cells grown under ALI conditions.^{39,48} Further studies characterizing the effects of SCFAs on TGF- β 1 and IL-9-induced barrier dysfunction will clarify the full impact of SCFA treatment on the compromised esophageal barrier.

We considered signaling via free fatty acid receptors GPR41, GPR43, and GPR109a as a potential mechanism for the barrier-restorative effects of butyrate and propionate. AR420626, 4-CMTB, and niacin, agonists for GPR41, GPR43, and GPR109a, were used to investigate whether the activation of these receptors could mimic the effects of SCFAs on EPC2. GPR agonists did not increase epithelial integrity as measured by TEER and FITC flux in IL-13-stimulated EPC2 ALI cultures contrasting the effects of butyrate and propionate. Also, *CAPN14* and *FLG* mRNA expression was unaffected by GPR agonists, indicating that the barrier-restorative effects of SCFAs are independent of GPR stimulation. Furthermore, as a positive control for GPR stimulation, we studied the effect of GPR agonists on LPS-induced IL-6 and IL-8 production by HUVECs, since it has been shown that this is partially mediated via GPRs.²¹ We observed a dose-dependent decrease in LPS-induced IL-6 and IL-8 production, indicating that the

lack of a response in EPC2 ALI cultures is not caused by biologically inactive GPR agonists but by the inability of these GPR agonists to induce SCFA-like effects. Our findings are in line with other studies demonstrating that SCFAs can exert their effects independent of free fatty acid receptors GPR41, GPR43, and GPR109a.⁴⁹⁻⁵¹

Alternatively, SCFAs can directly act as nuclear HDAC inhibitors.^{19,21,22} Indeed, both butyrate and propionate attenuated HDAC activity in EPC2. To further investigate whether HDAC inhibition could potentially contribute to the barrier-restorative effects of butyrate and propionate the pan-HDAC inhibitor TSA was used.^{21,49,50} TSA is structurally unrelated to butyrate and propionate but is 1000 times more potent in inhibiting HDAC than these SCFAs.⁵² HDAC inhibition results in histone hyperacetylation, leading to changes in chromatin structure that facilitate access for transcription factors to the promoter region of certain genes which then induces gene transcription. However, despite the overall correlation between histone acetylation and transcriptional activity, active gene transcription rather relates to the transcriptional competence of the gene than the high levels of histone acetylation.⁵³⁻⁵⁵ This could explain why the effects of TSA on barrier function in EPC2 ALI cultures measured by TEER and FITC flux were modest compared with those of butyrate and propionate. Thus, the ability of these SCFAs to directly inhibit HDAC activity may only be in part involved in their barrier-restorative effects.

Nevertheless, our studies have some limitations. Exposure to air in the ALI culture is essential for terminal epithelial differentiation and stratification. EPC2 ALI cultures were therefore treated with SCFAs in the basolateral compartment, but similar high concentrations of SCFAs may be difficult to reach systemically.⁵⁶ However, our data on apical SCFA treatment suggest that SCFA exposure from the apical side of the epithelium also supports the restoration of the esophageal epithelial barrier. Interestingly, it has been shown that increased dietary fiber intake influences the esophageal microbiome, which might lead to increased local SCFA concentrations in the esophagus.⁵⁷ Furthermore, we used the immortalized human esophageal epithelial cell line EPC2-hTERT to study the effects of SCFA treatment on an IL-13-compromised barrier. It may support our study to confirm our findings in primary human esophageal epithelial cells derived from EoE patients despite the marked transcriptional and morphologic overlap between IL-13-stimulated EPC2 ALI cultures and inflamed esophageal tissue.^{3,6}

The esophageal epithelial barrier during active EoE is impaired and selectively permeable to food allergens that can remain in the esophageal epithelium for up to 4 days.⁵⁸ The presence and subsequent recognition of food allergens in the esophageal mucosa generates a local type 2 immune response,⁵⁹⁻⁶¹ forming a pathogenic cycle to further exacerbate allergic inflammation. Butyrate and propionate may break this cycle by restoring barrier function and thus preventing the penetration of food allergens into the esophageal mucosa and subsequent inflammation.

The interest in dietary therapies for EoE has recently emerged as a result of the limitations associated with other therapies, and its effectiveness in achieving and maintaining clinical remission while

avoiding the need for drugs.⁶² A recent meta-analysis has shown that empiric elimination diets have moderate response rates (71%), but require a large number of endoscopies, whereas the efficacy of skin allergy testing-directed food elimination is questionable (45%).⁶³ Interestingly, complete dietary allergen avoidance using an elemental diet is highly effective in both children and adults (90.8%),⁶³ and restores esophageal mucosal integrity.^{44,64} It would be interesting to investigate whether a dietary intervention with SCFA formulations could restore esophageal immune fitness and improve symptoms.

In conclusion, our findings demonstrate that butyrate and propionate restore esophageal barrier function after IL-13-induced impairment, and that this is at least in part mediated by their ability to directly inhibit HDAC activity. Deeper knowledge of the mechanisms underlying the beneficial effects of butyrate and propionate could lead to novel approaches to restore esophageal barrier function. Our data highlight a potential role for butyrate and propionate in the management of EoE.

ACKNOWLEDGMENTS

This research is funded within the Partnership between NWO domain Applied and Engineered Sciences and Danone Nutricia Research and with additional financial support from Topsector Agri and Food, project number 16495 with the acronym LOIRE. We thank Dr. Anil Rustgi (University of Pennsylvania) for the EPC2-hTERT cell line.

CONFLICTS OF INTEREST

MTJVA, JG and BCAMVE are partly employed by Danone Nutricia Research and salary does not depend on the outcomes of this study. All other authors report no conflict of interest.

AUTHOR CONTRIBUTIONS

Study design: MTAK, FAR, and BCAMVE. Data collection: MTAK with assistance of BMH. Data analysis and interpretation: MTAK, FAR, and BCAMVE. Drafting of the manuscript: MTAK. Critical review of the manuscript: MLH, MTJVA, AJB, JG, FAR, and BCAMVE. All authors have read and agreed to the published version of the manuscript.

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SUPPORTING INFORMATION

Additional supporting information may be found online in the Supporting Information section.

How to cite this article: Kleuskens MTA, Haasnoot ML, Herpers BM, et al. Butyrate and propionate restore interleukin 13-compromised esophageal epithelial barrier function. *Allergy*. 2022;77:1510–1521. <https://doi.org/10.1111/all.15069>