Akkermansia muciniphila **identified as key strain to alleviate gut**

barrier injury through Wnt signaling pathway

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Credit author statement

XH and XM: Designed the experiments; XM and TX: Performed the experiments; XZ

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ABSTRACT

As the largest mucosal surface, the gut has built a physical, chemical, microbial and immune barrier to protect the body against pathogen invasion. The disturbance of gut microbiota aggravates pathogenic bacteria invasion and gut barrier injury. Fecal microbiota transplantation (FMT) is a promising treatment for microbiome-related disorders, where beneficial strain engraftment is a significant factor influencing FMT outcomes. The aim of this research was to explore the effect of FMT on antibiotic-induced microbiome-disordered (AIMD) model infected with enterotoxigenic *Escherichia coli* (ETEC). We used piglet, mouse and intestinal organoid models to explore the protective effects and mechanisms of FMT on ETEC infection. The results showed that FMT regulated gut microbiota and enhanced the protection of AIMD piglets against ETEC K88 challenge, as demonstrated by reduced intestinal pathogen colonization and alleviated gut barrier injury. *Akkermansia muciniphila* (*A. muciniphila*) and *Bacteroides fragilis* (*B. fragilis*) were identified as two strains that may play key roles in FMT. We further investigated the alleviatory effects of these two strains on ETEC infection in AIMD mice model, which revealed that *A. muciniphila* and *B. fragilis* relieved ETEC-induced intestinal inflammation by maintaining the proportion of Treg/Th17 cells and epithelial damage by moderately activating the Wnt/β-catenin signaling pathway, while the effect of *A. muciniphila* was better than *B. fragilis*. We therefore identified whether *A. muciniphila* protected against ETEC infection using basal-out and apical-out intestinal organoid models. *A. muciniphila* did protect the intestinal stem cells and stimulate proliferation and differentiation of intestinal epithelium, and the protective effects of *A. muciniphila* was reversed by Wnt inhibitor. FMT alleviated ETEC-induced gut barrier injury and intestinal inflammation in AIMD model. *A. muciniph*ila was identified as key strain in FMT to promote proliferation and differentiation of intestinal stem cells by mediating the Wnt/β-catenin signaling pathway.

Keywords: *Akkermansia muciniphila*; Wnt/β-catenin signaling pathway; Apical-out intestinal organoids; Gut barrier injury; Fecal microbiota transplantation.

Introduction

Growing evidence suggests that gut microbiota and its metabolites play important roles in modulating host health (*Canfora et al., 2019; Ghosh et al., 2021; Zheng et al., 2021*). The most drastic exposure that leads to intestinal microbiota dysbiosis is antibiotic treatment, which kills commensal micro-organisms and inhibits these microbiota-mediated immune defense (*Andremont et al., 2021; Buffie and Pamer, 2013*). The disturbance of intestinal microbiota may lead to disruptions of gut barrier and increase the susceptibility of host to pathogenic microbes (*Flint et al., 2012; Witkowski et al., 2020*). Several clinical studies have demonstrated that early-life exposure to antibiotic leads to gut microbiota dysbiosis that can impair host immune system maturation (*An et al., 2014; Nguyen et al., 2020*). Furthermore, the negative impact of antibiotics on gut microbiota in early life may even last for long periods throughout a lifetime, increasing the risk of gut diseases (*Francino, 2016*). Enterotoxigenic *Escherichia coli* (ETEC) K88 infection is a common cause of diarrhea in humans and animals (*Dubreuil, 2021*). When ETEC adheres to intestinal epithelial cells, virulence factors interact with specific receptors to induce intestinal barrier injury and intestinal inflammatory responses (*Yu et al., 2018*). Fecal microbiota transplantation (FMT) is a therapeutic intervention for reconstructing gut microbiota in gastrointestinal inflammatory diseases (*Borody et al., 2019; Colman and Rubin, 2014*). Since FMT has been successful in treating *Clostridioides difficile* infection and maintaining gut barrier, emerging investigation has been focused on other diseases (*Quraishi et al., 2017*). However, whether FMT could remedy against ETEC K88 infection in antibiotic-induced gut microbiota-disordered (AIMD) piglets is still unknown. Simultaneously, the underlying mechanism of FMT and the gut microbes that confer its efficacy remain unclear.

Intestinal stem cell renewal is vital for the maintenance of gut barrier (*Beumer and Clevers, 2021*). Mammalian intestinal epithelial cells are renewed approximately every 3-5 days with the migration of undifferentiated intestinal cells from the crypt to the top of the villi, accompanied by the differentiation of different types of epithelial cells (*Luo et al., 2022*). The proliferation and differentiation of intestinal epithelium driven by intestinal stem cells, mainly

crypt base columnar cells expressing R-Spondin receptor Lgr5, is a necessary process for

repairing intestinal barrier injury to prevent pathogen invasion (*Yan et al., 2017*). *Sato et al.*

(*2009*) successfully cultured isolated intestinal stem cells into three-dimensional intestinal

organoids with villous and crypt-like structural morphology for the first time. Intestinal

organoids contain various terminally differentiated cell types, including intestinal stem cell,

tuft cell, absorptive cell, enteroendocrine cell, goblet cell, and Paneth cell, which have more

advantages over single cell lines for *in vitro* study of intestinal regeneration (*Sprangers et al.,*

2021). Intestinal organoid is an ideal model for studying the interaction between intestinal

epithelium and microorganisms in vitro, which can eliminate the complexity of animal models

(*Yin et al., 2019*). In traditional basal-out intestinal organoid models, the intestinal epithelium

was in the interior of organoid spheroid, which restricted the interaction between intestinal

epithelium and gut microbiota. *Co et al.* (*2019*) first established the intestinal organoid model

with polarity reversal in human and mouse, which optimizes the shortcomings of

microinjection in terms of heterogeneous exposure concentrations and durations, as well as

destruction of organoid walls, and evaluated the infection effects of the invasive pathogens

with different polarity-specific patterns.

In the present study, we first investigated the effects of FMT on ETEC K88 infection in

AIMD neonatal piglets, then by using bacterial 16S rDNA sequencing analysis, we identified

Akkermansia muciniphila (*A. muciniphila*) as potential microbe to alleviate ETEC-induced

intestinal barrier injury. We further hypothesized that *A. muciniphila* could relieve intestinal

barrier injury and intestinal inflammation by modulating the proliferation and differentiation

of intestinal epithelium, and verified this hypothesis using *in vivo* mice experiments as well as

in vitro porcine intestinal organoid models. Collectively, our results may provide theoretical

basis that *A. muciniphila* is a promising method to repair intestinal barrier damage and a new

strategy for the precise application of *A. muciniphila* in livestock production.

Materials and methods

Ethics statement

All animal experiments were performed according to the guidelines of the Animal Care

Committee of Zhejiang University School of Medicine (permit number SYXK 2012-0178).

The animal care protocol was approved by the Local Committee of Animal Use following the

Guide for the Care and Use of Laboratory Animals (NIH publication 86-23 revised 1985).

Preparation of fecal microbiota suspension and bacterial strains

The fecal microbiota suspension was prepared according to our previous method (*Ma et*

al., 2021). In short, healthy adult Jinhua pigs that administrated medicinal feed additives or

drugs for more than 3 months were selected as fecal donors. The potential risks and infectious

pathogens in donor pigs were determined by serological and stool testing to ensure the safety

of the porcine FMT. Fresh fecal samples that collected from Jinhua pigs were transported in

AnaeroPack (Mitsubishi Gas Chemical, Japan) on ice to laboratory within 2 hours. The 50 g

fecal samples were homogenized in 250 mL sterile phosphate-buffered saline solution (PBS),

filtered through sterile stainless-steel strainer, and then centrifuged at 4000 rpm for 10 min.

The precipitate was resuspended in sterile PBS solution containing 10% sterile glycerol, and

the fecal microbiota suspension was stored at -80°C. All the above preparation was performed

in an anaerobic incubator (AW400SG anaerobic workstations; Electrotek, UK). All the

facilities and tools for the preparation of the fecal microbiota suspension were sterilized prior

to use. When processing the FMT, the fecal microbiota suspension was thawed in 37°C water

bath. ETEC K88 serotype O149:K91:K88ac was purchased from the China Institute of

Veterinary Drug Control (Beijing China), cultured in Luria-Bertani medium (Qingdao Hope

Bio-Technology Company, China), and incubated at 37□ in a shaker with 250 rpm overnight.

A. muciniphila ATCC BAA-835 and *Bacteroides fragilis* (*B. fragilis*) ATCC 25285 were

purchased from Guangdong Microbial Culture Collection Center (GDMCC, Guangzhou,

China). *A. muciniphila* was cultured in brain heart infusion media (OXOID, Thermo Fisher

Biochemicals Ltd., UK) supplemented with 0.1% mucin (Sigma-Aldrich, USA). *B. fragilis*

was cultured in trypticase soy broth (Qingdao Hope Bio-Technology Company, China)

supplemented with 5% defibrinated sheep blood (Qingdao Hope Bio-Technology Company,

China). These two anaerobic bacteria strains were incubated at 37□ in an anaerobic incubator

with 10% H_2, 10% CO_2, and 80% N_2 (AW400SG anaerobic workstations; Electrotek, UK).

The cultures were centrifuged for 10 min at a speed of 4000 rpm, and the pellets were then

resuspended in either aerobic or anaerobic PBS.

Animals

A total of 9 litters (9-11 piglets per litter) of Duroc × Landrace × Yorkshire ternary

hybrid neonatal piglets (aged 5 days) with identical birth dates and parities were obtained

from Anji Lvjiayuan Animal Husbandry Co. Ltd. (Zhejiang, China). Piglets were individually

housed in pens with appropriate environment of 24-26□ and 55-65% humidity. All newborn

piglets were fed with artificial milk substitutes, in which the nutrients reached the

requirements recommended by the National Research Council (NRC, 2012), and had *ad*

libitum access to water.

A total of 130 female C57BL/6 mice aged 5 weeks were obtained from Shanghai SLAC

Laboratory Animal, Co., Ltd. (SCXK (Zhejiang) 2017-0005; Shanghai, China). Mice were

housed in vinyl isolators in a room with 23 ± 1□ room temperature, 55-65% humidity and 12

h/12 h light/dark schedule. All mice had free access to water and food.

Experimental design

The piglets were randomly assigned to three groups (three litters per group): the control

group (Con group), the ETEC K88 + PBS group (EP group) and the ETEC K88 + FMT group

(EF group). The piglets in EP and EF groups were orally inoculated with 2 ml ampicillin (120

mg kg^{-1}) twice daily on day 1-3 of the experiment to disorder the intestinal resident

microbiota, then orally inoculated with ETEC K88 (1×10^9 CFU ml^{-1}) suspended in 2 ml

sterile PBS once daily on day 4-6 of the experiment. After ETEC K88 infection, the piglets in

EF group received 1.5 ml fecal microbiota suspension once daily by oral gavage for 6 days,

while the piglets in EP group received the same volume of sterile PBS. Meanwhile, the piglets

in Con group were orally inoculated with the same volume of sterile PBS throughout the

experiment. The doses of ampicillin and fecal microbiota suspension were based on a

previous study by Ma et al. [18]. Experimental design of piglets is shown in Figure 1A. The

weight of each piglet was measured at both the beginning and the end of the experiment to

calculate the average daily gain (ADG), and diarrhea score (0, normal feces; 1, moist feces; 2,

mild diarrhea; 3, severe diarrhea) was recorded throughout the experiment. Six piglets (2

piglets per litter) were picked from each group to be slaughtered at the end of the experiment

(Day 13).

After acclimating for a week, mice were randomly assigned to four groups (n = 10 each):

the control group (C group), the ETEC K88 + PBS group (E group), the ETEC K88 + *A.*

muciniphila group (A group) and the ETEC K88 + *B. fragilis* group (B group). On day 1-3 of

the experiment, the mice in E, A and B groups were orally inoculated with 0.2 ml 15 mg ml^{-1}

ampicillin (Sigma-Aldrich, USA) twice daily to disorder the gut microbiota. On day 4-6 of the

experiment, the mice in E, A and B groups were infected with ETEC K88 (1×10^9 CFU)

suspended in 0.2 ml 0.1 M $NaHCO_3$ buffer (pH 9.0) by oral gavage once daily. Mice were

fasted for 12 h before infection. Meanwhile, the mice in C group were given the equal volume

of sterile PBS on day 1-6 of the experiment. On day 7-13 of the experiment, live *A.*

muciniphila (1×10^6 CFU) and live *B. fragilis* (1×10^8 CFU) suspended in 0.2 ml anaerobic

sterile PBS were administered intragastrically once daily to the mice in A group and B group

respectively, while mice in C and E groups received the same volume of anaerobic sterile PBS.

The design of mouse experiment is shown in Figure 1B. Mice were orally infused with 0.2 ml

40 mg ml^{-1} 4 kDa fluorescein isothiocyanate-dextran (Sigma-Aldrich, USA) 4 hours before

sampling. At the end of the experiment (Day 14), six mice in each group were randomly

selected for ether anesthesia and neck dislocation.

184 Culture and treatment of porcine intestinal organoid

The cryovial of porcine intestinal organoid was kindly provided by Professor Li

Xiaoliang (Institute of Preventive Veterinary Medicine, College of Animal Sciences, Zhejiang

University, Hangzhou, China). The frozen organoids were thawed in a 37□-water bath for 2

min, and then added 1 ml of DMEM/F-12 (Gibco, USA) with 1% bovine serum albumin

(BSA, BioFroxx, Germany) solution. The contents of the cryovial were mixed and transferred

to the 15 ml conical tube containing 2 ml of DMEM/F-12 with 1% BSA solution, and then

centrifuged at $200 \times g$ for 5 min at 4□. The organoids were resuspended in equal volume of

complete IntestiCult[TM] Organoid Growth Medium (06010, StemCell[TM] Technologies, Canada)

and Matrigel Matrix (356231, Corning, USA). The 50 μl of the 500-crypt suspension was

plated in the center of each well of the pre-warmed 24-well plate (3524, Corning, USA). The

plate was placed at 37□ with 5% CO_2 for 10 min and then added 750 μl complete

IntestiCult[TM] Organoid Growth Medium containing 100 μg ml^{-1} penicillin/streptomycin

(Invitrogen, USA) to each well. The culture medium was fully exchanged three times per

week. For the treatment, the frozen organoids were passaged two times after thawing.

For generation of apical-out organoids, the procedure of reversing the polarity of

organoids was performed according to a previously published protocol by *Co et al.* (*2021*).

Briefly, after growing in Matrigel Matrix with growth medium for 7 days, the Matrigel-

embedded organoids were gently dissolved with 500 μl cold 5 mM ethylene diamine

tetraacetic acid (EDTA) in D-PBS and transferred to the 15 ml conical tube containing 10 ml

of cold 5 mM EDTA. The tubes were incubated on a rotating platform at 4□ for 1 h, and then

centrifuged at $200 \times g$ for 3 min at 4□. The organoids were resuspended in complete

IntestiCult[TM] Organoid Growth Medium containing 100 μg ml^{-1} penicillin/streptomycin and

transferred to ultra-low attachment 24-well plate (3473, Corning, USA). The suspended

organoids were cultured at 37□ with 5% CO_2 for 3 days.

To induce intestinal damage, the culture medium of basal-out and apical-out organoids

was changed to DMEM/F-12 containing ETEC K88 (10^6 CFU). After the invasion for 1 h, the

organoids were transferred to DMEM/F-12 containing 200 μg ml^{-1} penicillin/streptomycin for

1 h to kill residual extracellular bacteria. Then, the organoids were incubated in DMEM/F-12

containing *A. muciniphila* (10^5 CFU) for 1 h to detect the repair effect of *A. muciniphila*. In

addition, the organoids in WNT-C59 group were pretreated with the Wnt inhibitor Wnt-C59

(100 nM, AdooQ, China) for 24 h prior to damage. After co-cultivation, the organoids were

collected and resuspended in 4% paraformaldehyde (PFA, Servicebio, China) at 4□ or

RNAiso Plus (Takara, Japan) at -80□ for further analysis. For epithelial barrier integrity, the

apical-out organoids were resuspended in a solution of 4 kDa fluorescein isothiocyanate-

dextran (FD4) (2 mg ml^{-1}) diluted in DMEM/F-12 without phenol red. The design of

organoids' experiment is shown in Figure 1C.

Real-time quantitative PCR (RT-qPCR) analysis

Total RNA from intestinal tissue and organoids was extracted using TRIzol method

according to the precise instructions and cDNA was synthesized using PrimeScript™ RT

Master Mix (Takara, Japan). With cDNA as template, RT-qPCR was performed using TB

Green$^®$ *Premix Ex Taq*™ (Takara, Japan) on a LightCycler 480 System (Roche, Germany).

The real-time primers used in this study are listed in Table S1. The relative mRNA expression

of target genes was statistically analyzed based on the $2^{-\triangle\triangle Ct}$ method.

Bacterial load and intestinal permeability analyses

The bacterial load of ETEC K88 in piglets' jejunal and colonic mucosa was determined

by the absolute quantification. In brief, the gene of ETEC K88 (Genbank Accession:

M25302.1) was amplified by PCR (mK88-F: GGAATGGAAAGTTGGTACAGGTCTT,

mK88-R: CCGGCATAAGATGCATTCACTTTC), and the PCR product was analyzed by

agarose gel electrophoresis. The specific band was excised and recovered, connected with

pGM-T vector, and transformed into high-efficiency chemoreceptor DH-5α cells. After

culturing, the plasmid DNA was extracted and sequenced. The RT-qPCR analysis was

performed as the description of section 2.5. The copy number of samples was calculated by

the copy number of standard and the standard curve.

For the detection of ETEC K88 load in mice, fresh feces, jejunum and colon tissue, liver,

spleen and kidney were homogenized in sterile normal saline using a tissue homogenizer

(Jingxin Industrial Development Co., Ltd., China), then the homogenates were diluted 10-fold

continuously, and the dilutions were plated on MacConkey Agar (Qingdao Hope Bio-

Technology Company, China). After incubating in aerobic incubator at 37□ for 24 h, the

plates were counted three times. The results were presented as Lg CFU g^{-1} of samples.

For intestinal permeability assessment in mice, the serum fluorescence was detected

using a SpectraMax M5 plate reader (Molecular Devices, USA) at a 485 nm excitation

wavelength and a 535 nm emission wavelength. The standard curve was established from the

fluorescence of FD4 at concentrations of 0.005, 0.01, 0.02, 0.04, 0.08, 0.16 0.32, 0.64, 1.28,

2.56, 5.12 and 10.24 μg ml^{-1} to calculate the FD4 concentration.

For detection of epithelial barrier integrity in organoids, the apical-out organoids in the

FD4 solution were transferred to the slide and instantly imaged live by the laser scanning

confocal microscope IX81-FV1000 (Olympus, Japan) with FV10-ASW software (Olympus,

Japan).

Histology and morphology analyses

Jejunum and colon tissues were collected, fixed in 4% PFA, dehydrated with gradient

alcohol and embedded in paraffin. The tissue sections were stained with hematoxylin-eosin

(H&E) solution. The indexes related to the morphology of jejunum and colon were measured

by using Image Plus (v6.0). Scanning electron microscopy (SEM) and transmission electron

microscopy (TEM) were performed according to the procedure of the Bio-ultrastructure

Analysis Lab of the Analysis Center of Agrobiology and Environmental Sciences, Zhejiang

University. In brief, intestinal tissues were fixed in 2.5% glutaraldehyde, washed with PBS

three times, postfixed with 1% OsO4 for 1 h, then dehydrated with gradient alcohol (30%,

50%, 70%, 80%, 90%, 95% and 100%) for about 15 min at each step. For SEM, the samples

were dehydrated in Hitachi Model HCP-2 critical point dryer, coated with gold-palladium in

Hitachi Model E-1010 ion sputter, and observed in Hitachi Model SU-8010 SEM. For TEM,

the infiltrated samples were embedded in Spurr resin, sectioned in LEICA EM UC7 ultratome,

and sections were stained by uranyl acetate and alkaline lead citrate, then observed in Hitachi

Model H-7650 TEM.

Immunofluorescence analysis

Jejunum tissues were fixed in 4% PFA, dehydrated, paraffin embedded and sectioned.

After antigen retrieval and blocking, the sections were incubated in primary antibodies at 4□

overnight and secondary antibodies at room temperature for 50 min in the dark, followed by

staining with DAPI at room temperature for 10 min in the dark. The fixed organoids were

incubated in primary antibodies in blocking/permeabilization buffer at room temperature for 4

h, and then incubated in secondary antibodies together with DAPI and phalloidin (Servicebio,

China) in blocking/permeabilization buffer at room temperature for 2 h in the dark. The

fluorescence images were captured using a laser scanning confocal microscope IX81-FV1000

(Olympus, Japan) combined with FV10-ASW software (Olympus, Japan). The information of

primary and secondary antibodies is shown in Table S2.

Western blot analysis

Total protein was extracted by radioimmunoprecipitation assay lysis buffer and protein

concentration was determined by a BCA Assay Kit (Thermo Fisher Scientific, USA). Equal

protein amounts (60 μg) were electrophoresed on 10% sodium dodecyl sulphate-

polyacrylamide gel and the separated proteins were transferred onto the polyvinylidene

difluoride membranes (Millipore, USA) The membranes were blocked with Tris-buffered

saline with Tween (TBST) containing 5% BSA for 1 h at room temperature, and incubated

with the primary antibodies (Table S3) in TBST containing 3% BSA overnight at 4□. After

several times washes in TBST, the membranes were incubated with secondary antibodies

(Table S3) in TBST containing 3% BSA for 1 h at room temperature. The protein bands were

visualized using SuperSignal$^®$ West Dura Extended Duration Substrate (Thermo Fisher

Scientific, USA), and quantified by Image J software.

Microbiome sequencing analysis

Total DNA was extracted from intestinal contents using the E.Z.N.A. $^®$Stool DNA Kit

(D4015, Omega, Inc., USA) according to the manufacturer's instructions. The quality of DNA

was detected by agarose gel electrophoresis. Specific primers 341 F (5'-

CCTACGGGNGGCWGCAG-3') and 805 R (5'-GACTACHVGGGTATCTAATCC-3') were

used for bacterial PCR amplification of 16S rDNA V3-V4 hypervariable region. The PCR

products were purified using AMPure XT beads (Beckman Coulter Genomics, USA) and

quantified by Qubit (Invitrogen, USA). Sequencing libraries were generated using Ion Plus

Fragment Library Kit (Thermofisher, USA). The quantity of the amplicon library was

assessed on Agilent 2100 Bioanalyzer (Agilent, USA) and the Library Quantification Kit for

Illumina (Kapa Biosciences, USA). The libraries were sequenced on Illumina NovaSeq

PE250 platform. Paired-end reads were assigned to samples based on their unique barcode

and truncated by cutting off the barcode and primer sequence. Paired-end reads were merged

using FLASH. Quality filtering on the raw reads were performed under specific filtering

conditions to obtain the high-quality clean tags according to the fqtrim (v0.94). Chimeric

sequences were filtered using Vsearch software (v2.3.4). Denoising was conducted using

UNOISE3 to correct for sequencing errors. The sequences were clustered into Operational

Taxonomic Units (OTUs) with 97% identity using Uparse software (v7.0,1001) for all Clean

Reads of the sample. Species annotation of OTU sequences was performed using the Mothur

method with the SSUrRNA database of Silva 132, and multiple sequence alignment was

performed using MUSCLE (v3.8.31) software to obtain the phylogenetic relationships of all

OTUs sequences. The data of each sample were normalized using a standard of sequence

number corresponding to the sample with the least sequences. Subsequent analysis of alpha

diversity and beta diversity were all performed based on the output normalized data. Alpha

diversity is applied in analyzing complexity of species diversity for a sample through 4

indices, including Chao1, Observed species, Shannon, Simpson, and all these indices in our

samples were calculated with QIIME2. Beta diversity was calculated by QIIME2, and the

Anosim, principal coordinate analysis (PCoA) and non-metric multidimensional scaling

(NMDS) analyses were drawn implemented through R software (v3.5.2) with anosim,

WGCNA, stats, ggplot2 and vegan packages. Linear discriminant analysis (LDA) effect size

(LEfSe) package was used for LEfSe analysis. Other diagrams were implemented by using

the R software (v3.5.2).

Quantification of Short chain fatty acids

The colonic contents (50 mg) were homogenized with 0.5 ml dH_2O in ball mill and then

centrifuged. The supernatant 1 (0.3 ml) was homogenized with 0.5 ml dH_2O and then

centrifuged. The supernatant 2 (0.5 ml) combined with supernatant 1 were mixed with 0.8 ml

of 2-Methylvaleric acid as internal standard solution and 0.1 ml 50% H_2SO_4. After

centrifuging, the supernatant was transferred into fresh glass vial for Gas chromatography-

mass spectrometry analysis, which was performed using an Agilent 7890B gas chromatograph

system coupled with an Agilent 5977B mass spectrometer. Short chain fatty acids (SCFA)

were identified and quantified by the retention time and standard curves of the standard

solutions.

Flow cytometry (FC) analysis

Intestinal lamina propria cells were isolated by using Mouse Lamina Propria

Dissociation Kit (Miltenyi Biotec, Germany) according to the instruction. For cell surface

staining, the cells were incubated with FC block at 2-8□ for 15 min, then were stained with

Live/Dead Dye (FVS510, BD Pharmingen, USA) at room temperature for 15 min, followed

by washing with Stain Buffer. Cells were incubated with specific fluorescent antibodies

(Table S4) at 2-8□ for 30 min to label immune cells. For intracellular factor staining, cells

were fixed and permeabilized using Transcription Factor Fix/Prem Buffer at 4□ for 40 min,

washed with 1 × Perm/Wash Buffer, and stained with intracellular markers at 4□ for 30 min.

The stained cells were washed and resuspended in PBS, and assessed by the 12-color

FACSCelesta flow cytometer (Becton, Dickinson and Company, USA). The single cells were

first gated on SSC-A vs FSC-A and FSC-A vs FSC-H, and then CD45$^+$Live$^+$ cells were gated

on single cells. The CD45$^+$CD3$^-$ cells were gated on CD45$^+$Live$^+$ cells. The CD4$^+$ T cells

(CD3$^+$CD4$^+$ cells) were gated on CD45$^+$CD3$^-$ cells, Tregs (CD25$^+$Foxp3$^+$ cells), and Th17

cells (CD4$^+$RORγt$^+$ cells) were gated on CD3$^+$CD4$^+$ cells. The CD86$^+$CD11C$^+$ cells were

gated on DC$^+$CD3$^-$ cells. The data were analyzed using FlowJo software (v10.8.0).

Isolation and culture of mouse intestinal crypts

The jejunum proximal to the stomach was harvested immediately after mice were

sacrificed. After gently flushing with cold D-PBS (without calcium and magnesium) using

injection syringe, the intestinal segment was opened longitudinally and gently washed with

cold D-PBS three times. Then, the washed intestine was cut into 2 mm pieces with scissors,

which fell into a 50 ml conical tube containing 15 ml cold D-PBS. The rinsing procedure that

using the 10 ml serological pipette to gently pipette the intestinal pieces up and down three

times and then aspirate off the supernatant and add 15 ml fresh cold D-PBS was repeated 10-

15 times until the supernatant was clear. The intestinal pieces were resuspended in 25 ml

Gentle Cell Dissociation Reagent (GCDR, StemCell^{TM} Technologies, Canada) and incubated

at room temperature for 15 min on a rocking platform. After incubation, the intestinal pieces

were resuspended in 10 ml cold D-PBS containing 0.1% BSA, supernatant was filtered

through a 70 μm filter (Corning, USA), and filtrate was collected and labeled as fraction 1.

Repeat the above step three times to obtain fractions 2-4. The fractions were centrifuged at

290 × g for 5 min at 4□ and resuspended in 10 ml cold DMEM/F-12 with 15 mM HEPES.

The quality of the fractions was assessed by using an inverted microscope. The fraction which

enriched for intestinal crypts was selected, centrifuged and resuspended in equal volume of

complete IntestiCult^{TM} Organoid Growth Medium (06005, StemCell^{TM} Technologies, Canada)

and Matrigel Matrix. The 50 μl of the 500-crypt suspension was plated in the center of each

well of the pre-warmed 24-well plate. The plate was placed at 37□ with 5% CO_2 for 10 min

and then added 750 μl complete IntestiCult^{TM} Organoid Growth Medium containing 100 μg

ml^{-1} penicillin/streptomycin to each well. The culture medium was fully exchanged three

times per week. After 7 to 10 days of culture, the organoids were passaged using 1:6 split ratio.

The images of organoids were captured using inverted microscope (Nikon, Japan). The

surface areas of organoids were measured using ImageJ software (v1.8.0). The forming

efficiency (%) = (number of mature organoids growing after 5 days / number of crypts seeded)

× 100%.

Statistical analysis

Data were statistically analyzed by SPSS (v26.0) software, and all data are presented as

means ± standard deviation (SD). *P* value < 0.05 was considered significant. Kolmogorov-

Smirnov test was used to determine whether the data followed the normal distribution.

Comparisons between two groups were executed by unpaired Student's *t*-test or Mann-

Whitney *U*-test, and comparisons among three groups were executed by one-way ANOVA or

Kruskal-Wallis followed by Dunn's multiple comparisons.

Results

FMT enhanced the protection of AIMD piglets against ETEC K88 challenge

ETEC K88 infection is a leading cause of diarrhea in humans and animals. We set out to

explore whether FMT protected against ETEC K88 challenge. The piglets were randomly

assigned to three groups: the control piglets (Con group), the ETEC K88 infected piglets (EP

group) and the ETEC K88 infected piglets with FMT treatment (EF group). As shown in

Figure 2A and 2B, decreased ADG and increased diarrhea score were observed in AIMD

piglets at day 3 and day 6 after ETEC K88 infection, and these symptoms were relieved by

FMT treatment. We next detected the pathogen colonization in mucosal tissues and found that

FMT significantly decreased the copies of ETEC K88 in the jejunal and colonic mucosa

(Figure 2C). To address whether FMT could protect intestinal barrier against ETEC K88

infection, intestinal barrier integrity and inflammatory cytokine levels were detected. FMT

decreased the expression of proinflammatory cytokines (*TNF-α*, *IL-1β* and *IL-6*) and

increased levels of anti-inflammatory cytokines (*IL-10* and *TGF-β1*) in the jejunum (*P* < 0.01,

Figure 2D). Jejunal scanning electron microscopy showed that the villi (Figure 3A top) in

ETEC K88 infected piglets were short and coarse, and there existed obvious rod-shaped

bacterial adhesion on the surface of microvilli (Figure 3A bottom), while FMT reversed villi

and microvilli injury caused by ETEC K88. In addition, the reduced mRNA and protein levels

of the tight junctions (ZO-1, claudin and occludin) and adheren junctions (β-catenin and E-

cadherin) caused by ETEC K88 challenge were also reversed by FMT treatment in the

jejunum (Figure 3B-C). We then evaluated the expression of Mucin 2 (MUC2) in the jejunum

using immunofluorescence staining. As shown in Figure 3D, ETEC K88 infection reduced the

expression of MUC2, which was significantly increased following FMT administration. These

results indicated that FMT enhanced the protection of AIMD piglets against ETEC K88

challenge.

A. muciniphila **and** *B. fragilis* **were identified as two key strains in FMT**

We next assessed the variation of gut microbiota and SCFA metabolites in ETEC K88-

challenged piglets. The alpha diversity indexes were increased after FMT treatment (Figure

S1A-B). The PCoA and NMDS showed that gut microbiota between two groups clustered

separately (Figure S1C-D). The Anosim analysis (R = 0.396, *P* = 0.02) and UPGMA

clustering analysis also indicated that FMT altered the structure of the gut microbiota in

ETEC K88-challenged piglets (Figure 4A-B). LEfSe analysis verified that *B. fragilis* and

Akkermansia were changed significantly with an upper 4 LDA score after FMT administration

(Figure 4C). We compared the OTU sequence of *Akkermansia* in the National Center for

Biotechnology Information database by using Blast method based on Best Hit algorithm, and

the result was *A. muciniphila*. Furthermore, FMT significantly increased the production of

SCFA metabolites, including acetic acid, propionic acid, butyric acid, isobutyric acid,

isovaleric acid and hexanoic acid (Figure 4D). These results demonstrated that FMT regulated

gut microbiota and SCFA metabolites in ETEC K88-challenged piglets. *A. muciniphila* and *B.*

fragilis were identified as two strains that may play key roles in FMT.

A. muciniphila **and** *B. fragilis* **maintain intestinal barrier function of ETEC-induced**

mice

We further investigated whether these two strains exerted protective effects on ETEC

infection in AIMD mice model. Mice were randomly assigned to four groups: the control

mice (C group), the ETEC K88 infected mice (E group), the ETEC K88 infected mice treated

with *A. muciniphila* (A group) and the ETEC K88 infected mice treated with *B. fragilis* (B

group). As shown in Figure S2A and 5A, ETEC K88 infection led to weight loss and intestinal

morphology damage, which was relieved by *A. muciniphila* and *B. fragilis* treatment.

Correspondingly, scanning electron microscopy showed that *A. muciniphila* and *B. fragilis*

increased the height and number of microvilli and ameliorated the distribution of intercellular

junction in the jejunum and colon (Figure 5B). In addition, the relative mRNA and protein

expressions of MUC2 were significantly increased in the jejunum of *A. muciniphila* and *B.*

fragilis treated mice (Figure 5C). However, only *A. muciniphila* could upregulated MUC2

protein expression in the colon of treated mice (Figure 5C). Furthermore, we investigated the

diffusion of ETEC K88 in feces and organs after infection. ETEC K88 infection surged the

number of *Escherichia coli* in the feces, jejunum, colon, liver, spleen and kidney (Figure S2B).

A. muciniphila and *B. fragilis* treatment dramatically decreased the amount of ETEC K88 in

these tissues (Figure S2B), indicating a reduced translocation of ETEC K88 in the tissues and

organs. The concentration of FD4 in the serum has been commonly used as an indicator of

intestinal permeability. As shown in Figure S2C, *A. muciniphila* and *B. fragilis* significantly

decreased serum FD4 concentration, indicating *A. muciniphila* and *B. fragilis* enhanced

intestinal barrier integrity.

ETEC K88 adheres to intestinal epithelial cells and induces gut inflammation. As

demonstrated in Figure 6A, ETEC K88 decreased the relative mRNA expression of anti-

inflammatory cytokines *IL-10* and *TGF-β* and increased levels of pro-inflammatory cytokines

TNF-α, *IL-1β*, *IL-6* and *IFN-γ*, which was reversed in A and B groups, indicating that *A.*

muciniphila and *B. fragilis* ameliorated ETEC K88-induced intestinal inflammation. Intestinal

inflammatory response is closely related to immune cells. Treg cells secrete anti-inflammatory

cytokines IL-10 and TGF-β to alleviate intestinal inflammation. Conversely, Th17 cells

produce pro-inflammatory cytokines, such as IFN-γ and IL-17, which aggravate intestinal

inflammation. The dynamic balance of Treg and Th17 cells plays an important role in

intestinal immunity. Therefore, we used flow cytometry to analyze the proportion of the

immune cells isolated from jejunal lamina propria in the innate and adaptive immune system.

Treg cells were labeled with $CD25^+Foxp3^+$ cells, Th17 cells were labeled with $CD4^+ROR\gamma t^+$

cells, and mature dendritic cells were labeled with $CD86^+CD11C^+$ cells. ETEC K88 infection

significantly decreased the proportion of Treg cells and mature dendritic cells and increased

the proportion of Th17 cells, while oral administration of *A. muciniphila* or *B. fragilis*

significantly reduced the proportion of Th17 cells and increased the proportion of mature

dendritic cells (Figure 6B). Moreover, compared with E group, the proportion of Treg cells in

A group was significantly increased ($P < 0.01$), and the proportion of Treg cells in the B

group had an increasing trend ($P = 0.59$), indicating FMT maintained the proportion of

Treg/Th17 cells in ETEC K88-infected mice.

A. muciniphila **accelerated intestinal epithelial proliferation**

Since oral administration of *A. muciniphila* and *B. fragilis* mitigated intestinal mucosal

injury, we set out to explore their effects on the proliferation and differentiation of intestinal

epithelium. ETEC K88 infection prominently decreased *Ki67* and *Lyz* mRNA expression and

the number of $Ki67^+$ cells and Lyz^+ Paneth cells in the crypt of the jejunum (Figure 7A and

7B). *A. muciniphila* and *B. fragilis* increased *Ki67* and *Lyz* mRNA expression and the number

of $Ki67^+$ cells. However, only *A. muciniphila* enhanced the number of Lyz^+ Paneth cells,

suggesting that *A. muciniphila* accelerated the proliferation and differentiation of intestinal

epithelial cells. Lgr5-labeled active intestinal stem cells are the driving force of intestinal

epithelial proliferation and differentiation after intestinal injury. ETEC K88 infection

significantly increased the relative mRNA expression of *Lgr5*, *Wnt3*, *Axin2*, *Ctnnb1* in

jejunum mucosa and had no significant impact on *Notch1* and *Hes1* expression (Figure 7C). *A.*

muciniphila and *B. fragilis* inhibited the upregulation of *Lgr5*, *Wnt3*, *Axin2* and *Ctnnb1* at

transcriptional level caused by ETEC K88 infection and had no significant effect on *Notch1*

and *Hes1* levels. Western blot analysis also showed that *A. muciniphila* and *B. fragilis*

inhibited the upregulation Lgr5, Wnt3, Active β-catenin, c-Myc and CyclinD1 at protein level,

and the effect of *A. muciniphila* was better than that of *B. fragilis* (Figure 7D). These results

implied that *A. muciniphila* regulated the expression of intestinal stem cells in ETEC K88-

challenged mice and regulated Wnt/β-catenin signaling pathway rather than Notch signaling

pathway.

499   We next isolated jejunal crypt from mice and explore whether *A. muciniphila* or *B.*

fragilis influenced the ability of intestinal stem cells growing into intestinal organoids *ex vivo*.

In ETEC K88-infected mice, the crypts formed into undifferentiated spherical organoids,

while in *A. muciniphila* or *B. fragilis* treated mice, the crypts generated mature organoids with

bud structure (Figure 8A). We statistically analyzed the surface area and forming efficiency of

the intestinal organoids. As shown in Figure 8B, the organoids derived from ETEC K88-

infected mice had smaller surface area and lower forming efficiency than those from control

mice. *A. muciniphila* treatment led to larger surface area and higher forming efficiency of the

intestinal organoids. The surface area of intestinal organoids in B group was larger than that in

E group, while there was no significant difference in forming efficiency between these two

groups. Meanwhile, the relative mRNA expression of genes related to the proliferation and

differentiation at the intestinal organoid level was consistent with the results of jejunal

mucosa *in vivo* (Figure 8C). These results suggested that *A. muciniphila* accelerated intestinal

epithelial proliferation and regulated Wnt/β-catenin signaling pathway *ex vivo*.

***A. muciniphila* protected intestinal organoids against ETEC infection via Wnt signaling**

515   To explore the effects and mechanisms of *A. muciniphila* on ETEC K88 infection *in vitro*,

the porcine intestinal organoids were infected with ETEC K88 (ETEC group), and then co-

cultured with *A. muciniphila* (AKK group). Wnt inhibitor Wnt-C59 was added to explore

whether *A. muciniphila* acted through Wnt/β-catenin signaling pathway (WNT-C59 group),

and porcine intestinal organoids without any treatment were used as blank control (CON

group). The relative mRNA expression of *villin*, *ZO-1*, *Ki67*, *Lyz*, *MUC2*, *Lgr5*, *Wnt3a, β-*

catenin and the fluorescence intensity of villin, Ki67, Lgr5, Wnt3a and β-catenin in basal-out

intestinal organoids were significantly down-regulated upon ETEC exposure (Figure 9A-G).

A. muciniphila upregulated Lgr5, Wnt3a and β-catenin expression in ETEC-infected

organoids, while these effects were inhibited by Wnt-C59 treatment (Figure 9A-G). The

results suggested that *A. muciniphila* increased the number of intestinal stem cells and

activated Wnt signaling, but did not promote the proliferation and differentiation of basal-out

gut organoids. In basal-out intestinal organoid models, the intestinal epithelium was in the

interior of organoid spheroid, which restricted the interaction between intestinal epithelium

and *A. muciniphila*. Therefore, we reversed the polarity of basal-out intestinal organoids by

removing the Matrigel from the culture system and suspending intestinal organoids in the

culture (Figure S3A). In apical-out intestinal organoids, the apical surface was outward to

make it accessible to *A. muciniphila*. Under confocal microscopy, the F-actin of basal-out

intestinal organoid was located in the inside of the spheroids, while the F-actin of the apical-

out intestinal organoid was outside the spheroids, indicating that the polar reversal intestinal

organoid model was successfully constructed (Figure S3B). We conducted FD4 permeation

test in apical-out intestinal organoid model, and co-cultivation with *A. muciniphila* restored

the epithelial barrier integrity of organoids challenged by ETEC K88 (Figure 10A, H). *A.*

muciniphila not only increased the mRNA expression of *ZO-1*, *Lyz* and *MUC2* in the ETEC

K88 infected apical-out intestinal organoids, but also up-regulated the protein expression of

villin, Wnt3a, β-catenin and the number of Lgr5^{+} and Ki67^{+} cells (Figure 10B-G, I). Moreover,

A. muciniphila also relieved the inflammatory response of the ETEC K88 infected apical-out

intestinal organoids, as determined by decreasing proinflammatory cytokines, inducing *TNF-α*,

IL-1β, *IL-6* and *IFN-γ*, and increasing anti-inflammatory cytokines *IL-10* and *TGF-β* (Figure

10J). However, the moderating effect of *A. muciniphila* on the apical-out intestinal organoids

disappeared after Wnt-C59 exposure. Taken together, *A. muciniphila* accelerated the

proliferation and differentiation of the intestinal stem cells and ameliorated the intestinal

barrier injury and inflammation induced by ETEC K88. *A. muciniphila* protected intestinal

organoids against ETEC infection via Wnt signaling.

Discussion

The homeostasis of gut microbiota is a major contributor to host health. Intestinal

dysbiosis increases intestinal colonization of antibiotic-resistant and pathogenic bacteria.

Antibiotic treatment is the most drastic exposure leading to gut dysbiosis and pathogen

invasion (*Andremont et al., 2021*). Early-life exposure to antibiotic leads to gut microbiota

dysbiosis and impair host immune system maturation, which may even last for long periods

throughout life (*Nguyen et al., 2020*). Researchers have made a lot of attempts to prevent or

inhibit antibiotic-induced dysbiosis. Our previous study showed that FMT alleviates early-

life antibiotic-induced gut microbiota dysbiosis and mucosa injuries in a neonatal piglet

model (*Ma et al., 2021*). Our present study aimed to investigate whether FMT increases early-

life antibiotic induced neonatal piglet against pathogen invasion. ETEC K88 is one of the

most common pathogens in humans and animals and causes intestinal inflammation and

diarrhea symptoms (*Sun and Kim et al., 2017*). Studies demonstrated that ETEC K88

infection led to pro-inflammatory cytokine upregulation and intestinal mucosal barrier injury

(*Wang et al., 2020; Xie et al., 2021*). In ETEC-induced neonatal piglets, ZO-1 and Claudin-2

levels in ileum of were significantly decreased after ETEC K88 infection (*Xie et al., 2021*). In

accordance with the above results, the present study showed that ETEC K88 infection

increased the expression of pro-inflammatory cytokines and reduced the expression of

intercellular junction proteins in AIMD piglets. Since FMT has been successful in treating

Clostridioides difficile infection and maintaining gut barrier, emerging investigation has been

focused on other diseases (*Cammarota et al., 2017*). *Geng et al.* (*2018*) demonstrated that

FMT diminished the inflammatory response and the destruction of epithelial integrity in

piglets challenged with lipopolysaccharide. In the present study, we found that FMT

dampened ETEC K88-induced pro-inflammatory cytokines upregulation and improved the

intestinal morphology. In addition, the number of ETEC K88 in jejunal and colonic mucosa

was significantly decreased following FMT, which was consistent with the observation of

jejunal microvilli under SEM, showing the absence of rod-shaped bacterial adhesion in FMT

treatment. ETEC infection was reported to reduce the MUC2 expression by destroying goblet

cells or directly degrading MUC2, thereby damaging the integrity of the intestinal epithelial

barrier (*Luo et al., 2014*). Our results demonstrated that FMT enhanced the expression of

MUC2 and increased the expression of ZO-1, claudin, occludin, β-catenin and E-cadherin at

mRNA and protein levels, indicating that FMT improved the intercellular junctions between

intestinal epithelial cells and restored the integrity of the intestinal mucosal barrier to reduce

the colonization and invasion of ETEC K88.

Intestinal microbiota plays a key role in maintaining gut mucosal barrier and resistance

to pathogen invasion (*Chelakkot et al., 2018*). Antibiotic-induced gut microbiota-disorders

have been reported to disrupt gut barrier and increase the susceptibility of host to pathogenic

microbes (*Wlodarska et al., 2011*). So, we further probed the changes of gut microbiota in

AIMD piglets infected with ETEC K88. The results showed that FMT regulated gut

microbiota and SCFAs metabolism in ETEC K88-challenged piglets. LEfSe analysis

manifested that *B. fragilis* and *A. muciniphila* were the key differential bacteria between three

groups. *A. muciniphila* utilizes mucin as the only carbon and nitrogen source for its growth in

the gut, and its metabolite SCFAs can facilitate MUC2 secretion by goblet cells to maintain

the dynamic balance of intestinal mucus layer (*Ottman et al., 2017*). *A. muciniphila*

negatively correlated with many diseases, including diabetes, obesity and inflammatory bowel

diseases (*Belzer C and de Vos, 2012; Castro-Mejía et al., 2016; Plovier et al., 2017*). It has

been reported that *A. muciniphila* effectively inhibited intestinal inflammatory response and

elevated epithelial barrier function (*Everard et al., 2013; Li et al., 2016; Zhang et al., 2018*).

Meanwhile, *Bacteroides fragilis* is one of the symbiotic anaerobes within the mammalian gut

and is also an opportunistic pathogen which often isolated from clinical specimens (*Sun et al.,*

2019). Although it was initially thought to be pathogenic, in the long-term evolution process,

Bacteroides fragilis colonized in the gut has established a friendly relationship with the host,

which is an essential component for maintaining the health of the host, especially for obesity,

diabetes and immune deficiency diseases (*Troy and Kasper, 2010*). *B. fragilis* could

decompose mucin by lipid-anchored enzymes and interact with intestinal epithelial cells,

thereby altering intestinal permeability and repairing intestinal epithelial barrier (*Donaldson*

et al., 2016; Hsiao et al., 2013). *Deng et al.* (*2016*) reported that *B. fragilis* altered

macrophage phenotype and increased the phagocytosis of macrophages to restrain pathogens

colonization. Hence, we speculated that FMT increased the relative abundance of *A.*

muciniphila and *B. fragilis* in the intestine and alleviate intestinal barrier injury and intestinal

inflammation caused by ETEC K88 infection.

Since *A. muciniphila* and *B. fragilis* were identified as two strains that may play key

roles in FMT, we further investigate whether *A. muciniphila* and *B. fragilis* exerted protective

effect on ETEC K88 infection in AIMD mice model. We found that *A. muciniphila* and *B.*

fragilis maintained normal gut morphology after ETEC K88 infection, repaired the integrity

of intestinal mucosal barrier as evidenced by the reduced FD4 concentration in serum,

colonized in the intestine and inhibit the translocation of ETEC K88, and reduced the

intestinal inflammation, which may have been attributed to the increased proportion of Treg

cells and mature dendritic cells and the decreased proportion of Th17 cells. These results are

consistent with the protective role of *A. muciniphila* and *B. fragilis* in dextran sulphate sodium

salt induced enteritis model (*Bian et al., 2019; Wang et al., 2021; Zhai et al., 2019*). Intestinal

stem cell renewal is vital to maintaining the gut barrier. Ki67 is widely expressed in

proliferating cells and presents a reliable indicator of cell proliferation activity (*Zheng et al.,*

2010). Ki67 reaches its peak during the division stage with short half-life period and easy

degradation (*Zheng et al., 2010*). In our study, *A. muciniphila* and *B. fragilis* increased the

number of Ki67^{+} cells and the mRNA expression of *Lyz*, suggesting that *A. muciniphila* and *B.*

fragilis accelerated the proliferation and differentiation of intestinal epithelium. Paneth cells

not only produce antimicrobial peptides and express key niche signals that regulate stem cells,

but also secrete activators that dynamically modulate Wnt/β-catenin signaling pathway (*Hou*

et al., 2020). The Wnt/β-catenin signaling pathway is essential for the proliferation and

differentiation of intestinal stem cells (*Ring et al., 2014*). When the Wnt receptors are

activated, the multiprotein degradation complex is rapidly inhibited, which allows β-catenin

to accumulate and subsequently migrate to the nucleus and binds to the transcription factors

of T cell factor family and lymphoid enhancer-binding factor family to drive the expression of

genes that regulate the stem cells, including Lgr5, c-Myc and Cyclin D1 (*Niehrs, 2012*). The

c-Myc and Cyclin D1 are cyclins that positively regulate cell cycle from the G1 phase to S

phase, thus inducing cell cycle progression and promote cell proliferation (*Shi et al., 2015*).

Zhu et al. (*2020*) reported that *A. muciniphila* restored intestinal mucosal damage with the

accelerated proliferation of intestinal epithelium through Wnt/β-catenin signaling pathway,

thus relieving infection caused by *Salmonella pullorum* in chicks. In the present study, *A.*

muciniphila moderately activated Wnt/β-catenin signaling pathway and maintain the suitable

expression of Wnt, Active-β-catenin, Lgr5, c-Myc and Cyclin D1. Consistent with the *in vivo*

results, *ex vivo* results of mouse intestinal organoids also demonstrated that *A. muciniphila*

repaired the damaged intestinal epithelium caused by the ETEC K88 infection by modulating

the Wnt/β-catenin signaling pathway. Simultaneously, *B. fragilis* also avoid the overactivation

of the Wnt/β-catenin signaling pathway caused by the ETEC K88 infection, while the impact

of *B. fragilis* was not as obvious as *A. muciniphila*.

Intestinal organoid is an ideal model for studying the interaction between intestinal

epithelium and microorganisms in vitro, which can eliminate the complexity of animal models

and avoid the unicity of the traditional intestinal epithelial cells (*Yin et al., 2019*). In order to

further verify whether *A. muciniphila* attenuated ETEC K88 infection through Wnt/β-catenin

signaling pathway, basal-out and apical-out porcine intestinal organoid models was used.

Notably, we found that the ETEC K88 infection down-regulated Wnt/β-catenin signaling

pathway *in vitro*, which was inconsistent with the results *in vivo*. A recent study showed that

the addition of 400 ng ml^{-1} STp, an enterotoxin of ETEC, could suppress Wnt/β-catenin

signaling pathway of porcine intestinal organoids (*Zhou et al., 2021*). It is speculated that the

reason for the difference results might be related to the difference in the processing time of the

in vivo and *in vitro* treatment. In spite of the slight stimulation of *A. muciniphila* on the

proliferation and differentiation of infected intestinal organoids, *A. muciniphila* up-regulated

the Wnt/β-catenin signaling pathway and had protective effects on intestinal stem cells. In

basal-out intestinal organoid models, the intestinal epithelium was in the interior of organoid

spheroid, which restricted the interaction between intestinal epithelium and *A. muciniphila*.

Co et al. (*2019*) first established the intestinal organoid model with polarity reversal in human

and mouse and evaluated the infection effects of the invasive pathogens with different

polarity-specific patterns, *Salmonella enterica serovar Typhimurium* and *Listeria*

monocytogenes, on polarity reversal intestinal organoids. Subsequently, *Li et al.* (*2020*) first

applied the porcine intestinal organoids with polarity reversal to investigate the infectivity and

antiviral immune responses of porcine transmissible gastroenteritis virus. In addition, studies

on host-pathogen interactions using polarity reversal intestinal organoids have also been

reported in the sheep and chickens (). But up to now, there are few reports on the interaction

between host and symbiotic microorganism using polarity reversal intestinal organoids. Here,

we utilized porcine polarity reversal intestinal organoids to make *A. muciniphila* directly

interacting with the intestinal epithelium. Our results showed that *A. muciniphila* alleviated

the injury of polarity reversal intestinal organoids induced by ETEC K88. In agreement with

the *in vivo* results, *A. muciniphila* restrain the intestinal inflammation caused by the ETEC

K88 infection in polarity reversal intestinal organoids. Additionally, the alleviatory effect of *A.*

muciniphila on ETEC K88 infection via the Wnt/β-catenin signaling pathway was confirmed

with the addition of Wnt inhibitor Wnt-C59. The promoted proliferation and differentiation in

the basal-out and apical-out intestinal organoids after co-culturing with *A. muciniphila* were

reversed with Wnt-C59.

In conclusion, we found that FMT relieved intestinal inflammation and improved

intestinal barrier function in AIMD piglets infected with ETEC K88. *A. muciniphila* was

identified as key strain in FMT to promote the proliferation and differentiation of intestinal

stem cells by mediating the Wnt/β catenin signaling pathway. The study revealed the possible

mechanism that FMT alleviates the intestinal barrier injury.

DECLARATIONS

Ethics approval and Consent to participate

All animal experiments were performed according to the guidelines of the Animal Care

Committee of Zhejiang University School of Medicine (permit number SYXK 2012-0178).

The animal care protocol was approved by the Local Committee of Animal Use following the

Guide for the Care and Use of Laboratory Animals (NIH publication 86-23 revised 1985).

Consent for publication

Not applicable.

Availability of data and material

The raw sequences derived from microbiome analysis for this study can be found in the

Sequence Read Archive (SRA) (https://www.ncbi.nlm.nih.gov/sra/PRJNA837047). And all

datasets analyzed in the present study are available from the corresponding author on

reasonable request.

Competing interests

The authors declared that the work was carried out in the absence of any commercial or

financial relationships that could be construed as a potential conflict of interest.

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 Figure legends:

Figure 1. Experimental designs of the present study. (A) Schematic diagram of the FMT administered to
the AIMD piglets infected with ETEC K88. (B) Schematic diagram of the alleviatory effects of *A.*
muciniphila and *B. fragilis* on ETEC K88 infection in mice. (C) Schematic diagram of the alleviatory
effects of *A. muciniphila* on ETEC K88 infection in porcine basal-out and apical-out intestinal organoids.

Figure 2. FMT improved the growth performance and inflammatory response of AIMD piglets
infected with ETEC K88. (A) Average daily weight gain (ADG) in the Con, EP and EF groups. (B)
Diarrhea score at days 3 and 6 post-infection. (C) The colonization of ETEC K88 in the jejunum and colon.
The melting curve demonstrated that only the ETEC K88 could be amplified by the primers we used. (D)
The relative mRNA expression of cytokines in the jejunum. Con: control group, EP: ETEC K88 + PBS
group, EF: ETEC K88 + FMT group. Data are expressed as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$, ***,
$P < 0.001$.

Figure 3. FMT improved the intestinal morphology and barrier function in AIMD piglets infected
with ETEC K88. (A) SEM images of villi (top) and microvilli (bottom) in the jejunum (scale bars = 300 or
2 μm). (B) The relative mRNA expression of tight junction proteins (ZO-1, claudin and occludin) and
adheren junctions (β-catenin and E-cadherin) in the jejunum. (C) The relative protein expression of tight
junction proteins (ZO-1, claudin and occludin) and adheren junction proteins (β-catenin and E-cadherin) in
the jejunum. (D) Immunofluorescence images of Mucin 2 (MUC2, green) in the jejunum (scale bars = 50
μm). Con: control group, EP: ETEC K88 + PBS group, EF: ETEC K88 + FMT group. Data are expressed
as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$. Data shown are representative of at least two independent
experiments.

Figure 4. FMT changed the composition of gut microbiota and the concentration of SCFAs in AIMD
piglets infected with ETEC K88. (A) Anosim analysis. (B) UPGMA clustering analysis based on
Unweighted unifrac distance. (C) LEfSe analysis with LDA score > 4. (D) Quantification of SCFA
metabolites (acetic acid, propionic acid, butyric acid, isobutyric acid, isovaleric acid and hexanoic acid). EP:
ETEC K88 + PBS group, EF: ETEC K88 + FMT group. Data are expressed as the mean ± SD. *, $P < 0.05$,
**, $P < 0.01$. Data shown are representative of at least two independent experiments.

Figrue 5. *A. muciniphila* and *B. fragilis* improved the intestinal morphology of mice infected with
ETEC K88. (A) H&E-stained images of jejunum (scale bars = 50 μm) and the histograms of villus height,
crypt depth and the ratio of villus height to crypt depth. (B) TEM images of jejunum and colon (scale bars
= 0.5 μm). (C) Immunofluorescence images of Mucin 2 (MUC2, green) in the jejunum and colon (scale
bars = 50 μm). C: control group, E: ETEC K88 + PBS group, A: ETEC K88 + *A. muciniphila* group, B:
ETEC K88 + *B. fragilis* group. Data are expressed as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$, ***, $P <$
0.001. Data shown are representative of at least two independent experiments.

Figure 6. *A. muciniphila* and *B. fragilis* regulated the expression of inflammatory cytokines and the
balance of Treg and Th17 cells in mice infected with ETEC K88. (A) The relative mRNA expression of
cytokines in the jejunum. (B) Flow cytometric dot plots and proportions of Treg (CD25⁺Foxp3⁺), Th17
(CD4⁺RORγt⁺) and mature dendritic cells (CD86⁺CD11C⁺) in jejunal lamina propria. C: control group, E:
ETEC K88 + PBS group, A: ETEC K88 + *A. muciniphila* group, B: ETEC K88 + *B. fragilis* group. Data
are expressed as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$, ***, $P < 0.001$. Data shown are representative of
at least two independent experiments.

Figure 7. *A. muciniphila* and *B. fragilis* enhanced the number of Ki67 cells and Lyz Paneth cells in
jejunal crypt of mice infected with ETEC K88 by moderately activating the Wnt/β-catenin pathway.

(A) The mRNA expression and immunofluorescence staining images of intestinal epithelial cell proliferation marker Ki67 (red). (B) The mRNA expression and immunofluorescence staining images of Paneth marker Lyz (green). (C) The relative mRNA expression of *Lgr5*, *Wnt3*, *Notch1* and other genes in jejunum mucosa. (D) The relative protein expression of Wnt/β-catenin pathway and its target genes in jejunum mucosa. C: control group, E: ETEC K88 + PBS group, A: ETEC K88 + *A. muciniphila* group, B: ETEC K88 + *B. fragilis* group. Data are expressed as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$, ***, $P < 0.001$. Data shown are representative of at least two independent experiments.

Figure 8. *A. muciniphila* **and** *B. fragilis* **promoted the activity of Lgr5$^+$ intestinal stem cells as proved by the growth of intestinal organoids.** (A) Morphology of *ex vivo* culture of crypts isolated from the jejunum of mice in the C, E, A and B groups (Day 5; 4 ×, scale bars = 200 μm; 20 ×, scale bars = 50 μm). (B) The surface area and forming efficiency of intestinal organoids. (C) The relative mRNA expression of genes related to the proliferation and differentiation of the intestinal organoids. C: control group, E: ETEC K88 + PBS group, A: ETEC K88 + *A. muciniphila* group, B: ETEC K88 + *B. fragilis* group. Data are expressed as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$, ***, $P < 0.001$. Data shown are representative of at least two independent experiments.

Figure 9. *A. muciniphila* **protected the Lgr5$^+$ intestinal stem cell and activated the Wnt/β-catenin signaling pathway of the ETEC K88-induced basal-out intestinal organoids.** (A) The relative mRNA expression of *villin*, *ZO-1*, *Ki67*, *Lyz1*, *MUC2*, *Lgr5*, *Wnt3a* and *β-catenin* genes. (B)-(F) Immunofluorescence images of villin (scale bars = 50 μm), Ki67 (scale bars = 50 μm), Lgr5 (scale bars = 20 μm), Wnt3a (scale bars = 50 μm) and β-catenin (scale bars = 50 μm). (G) Fold change of the mean fluorescence intensity. CON: control group, ETEC: ETEC K88 + DMEM/F-12 group, AKK: ETEC K88 + *A. muciniphila* group, WNT-C59: Wnt-C59 + ETEC K88 + *A. muciniphila* group. Data are expressed as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$, ***, $P < 0.001$. Data shown are representative of at least two independent experiments.

Figure 10. *A. muciniphila* **alleviated the intestinal epithelial injury of the ETEC K88-induced apical-out intestinal organoids.** (A) Confocal microscope visualization of the apical-out intestinal organoids incubated in FD4 solution (scale bars = 50 μm). (B) The relative mRNA expression of *villin*, *ZO-1*, *Ki67*, *Lyz1*, *MUC2*, *Lgr5*, *Wnt3a* and *β-catenin* genes. (C)-(G) Immunofluorescence images of villin, Ki67, Lgr5, Wnt3a and β-catenin (scale bars = 20 μm). (H) Quantification of the apical-out intestinal organoids incubated in FD4 solution that have intact barrier integrity. (I) Fold change of the mean fluorescence intensity. (J) The relative mRNA expression of cytokines in the apical-out intestinal organoids. CON: control group, ETEC: ETEC K88 + DMEM/F-12 group, AKK: ETEC K88 + *A. muciniphila* group, WNT-C59: Wnt-C59 + ETEC K88 + *A. muciniphila* group. Data are expressed as the mean ± SD. *, $P < 0.05$, **, $P < 0.01$, ***, $P < 0.001$. Data shown are representative of at least two independent experiments.