Bacillus coagulans BACO-17 Alone or in Combination with Galacto-Oligosaccharide Ameliorates Salmonella-Induced Diarrhea and Intestinal Inflammation

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Abstract: In this study, a diarrhea model was established by exposing rats to appropriate antibiotics and Salmonella. After an in vitro screening of prebiotics, fructo-oligosaccharide and galacto-oligosaccharide (GOS) were selected; their symbiotic potential and ability to ameliorate diarrhea symptoms and intestinal inflammation with Bacillus coagulans BACO-17 were evaluated in vivo. After a 27-day feeding experiment including antibiotic intervention and Salmonella infection, it was found that using B. coagulans BACO-17 alone and in combination with GOS as a synbiotic could render a better recovery by lowering diarrhea indexes by 26.9% and 18.7%, respectively. Compared with the negative control, the administration of this synbiotic mixture resulted in the most significant increase in fecal concentrations of total short-chain fatty acids (about 2-fold higher), with a promising improvement in disrupted gut microbial balance. It was worth noting that the administration of B. coagulans BACO-17 alone or in combination with GOS effectively reduced intestinal inflammation (27–31%) and mucosal necrosis (82%) over the negative control. These results suggested that B. coagulans BACO-17 and GOS could be exploited as a promising synbiotic mixture to relieve intestinal inflammatory diseases and improve gut health.

Keywords: Bacillus coagulans BACO-17; galacto-oligosaccharide; diarrhea; antibiotic; intestinal inflammation

1. Introduction

Antibiotic-associated diarrhea (AAD) is a common complication developed from systemic antibiotic treatment. Antibiotic therapy is a vital tool to kill pathogenic bacteria but can also potentially lead to the disturbance of indigenous microbiota in the intestine. Specifically, disruption of the normal balance of intestinal flora results in alterations in carbohydrates and bile acids metabolism, reduction in short-chain fatty acids production, and the emergence of enteric pathogens such as Salmonella [1].

Bacillus coagulans is an endospore-forming bacteria widely used as a probiotic agent. It can develop spores and possesses good stability against harsh processing conditions such as low pH and high temperature [2,3]. Following spore germination intragastically, B. coagulans vegetative cells can proliferate in the intestine, elevate short-chain fatty acid concentrations, and subsequently improve the intestinal milieu [4]. Previous research has substantiated different therapeutic activities of B coagulans, including attenuated gut dysbiosis, decreased occurrence of abdominal pain and bloating, and alleviation of diarrhea-predominant irritable bowel syndrome through supplementation of B. coagulans spores [5–7].
Co-supplementation with a prebiotic could enhance the potential probiotic effects of *B. coagulans* by elevating its colonization and survival rates [8]. As previously reported by van der Beek et al. [9], orally administrated prebiotics could be fermented in the large intestine to produce short-chain fatty acids (SCFAs) and exert beneficial effects on intestinal health. Another study also indicated that consumption of *B. coagulans* and prebiotics effectively elevates organic acid production and populations of beneficial bacteria in the large intestine [10].

In light of the AAD resulting from disruption of the commensal gut microbiota, it was believed that supplementing the diet with *B. coagulans* and prebiotics could be a reasonable approach to ameliorating AAD by modulating and restoring gut microbiota [11]. This present study aimed to compare the prebiotic potential of four different carbohydrate components, including xylo-oligosaccharide, fructo-oligosaccharide, galacto-oligosaccharide, and citrus pectin. The synbiotic effects of *B. coagulans* BACO-17 and selected prebiotics in a rat diarrhea model were established by challenging the animals with ampicillin and clindamycin together with *Salmonella enterica* evaluated. More specifically, changes in different fecal parameters (e.g., diarrhea index (DI), moisture, and SCFAs) and histological assessment of colonic tissue after the consumption of *B. coagulans* BACO-17 and selected prebiotics are discussed.

2. Materials and Methods

2.1. In Vitro Screening of Prebiotics

A pure strain of *B. coagulans* BACO-17 in spore form was provided by Syngen Biotech Co., Ltd. (Taipei City, Taiwan) and stored at 4 °C until used. In order to select appropriate prebiotics for *B. coagulans* BACO-17 to create a synbiotic preparation, in vitro growth of *B. coagulans* BACO-17 in the presence of four different carbohydrate components was analyzed and compared to the probiotic bacteria growth in the presence of glucose as a control group.

*B. coagulans* BACO-17 cell suspension (about 4.9 Log CFU/mL) was added as 10% (v/v) inoculum into the test tubes containing 9 mL of medium, in which glucose was substituted with each prebiotic (0.5%, w/v) in equal amounts separately. The prebiotics used in the limiting carbon source replacement included citrus pectin, fructo-oligosaccharide (FOS), galacto-oligosaccharide (GOS), and xylo-oligosaccharide (XOS). These prebiotics were obtained from Syngen Biotech Co., Ltd. (Taipei City, Taiwan). GYE culture medium was used as the substrate. *B. coagulans* BACO-17 was cultivated at 55 °C for 24 h, and its growth profile in different carbon sources and standard glucose-containing media was observed. The results were used as a reference for subsequent in vivo experiments.

2.2. Diets and In Vivo Experimental Design

The animal study protocol was approved by the Animal Care and Use Committee of National Chung Hsing University, and the laboratory animals were handled in accordance with the institutional ethical guideline.

In this experiment, a total of fifty-four 7-week-old Sprague Dawley (SD) rats weighing 239.3 ± 2.9 g were purchased from BioLASCO Company, Taiwan (IACUC approval number: 109–105). These animals were individually housed in stainless steel cages and placed in a room maintained at 22 ± 1 °C with a 12-h light/dark cycle. After one week of acclimation, SD rats were randomly divided into seven groups, including two control groups and five sample groups. During the whole experimental period, the five sample groups included one probiotic group, administrated with *B. coagulans* BACO-17 (BC group, 9 log CFU/day), two prebiotic groups, administrated with FOS (FOS group, 1.03 g FOS/kg bw/day) and GOS (GOS group, 0.83 g GOS/kg bw/day), and two synbiotic groups, administrated with *B. coagulans* BACO-17 combined with FOS (FB group) and GOS (GB group). The FB group was fed with a mixture of *B. coagulans* BACO-17 (9 log CFU) and FOS (1.03 g/kg bw) each day. The GB group was fed with a mixture of *B. coagulans* BACO-17 (9 log CFU) and GOS (1.03 g/kg bw) each day. The negative control group was fed a basal chow diet and treated
with the same diarrheal disease model as the sample groups. The normal control group was only provided with a basal chow diet (LabDiet 5001).

During the experiment, all rats had free access to LabDiet 5001 and drinking water. Samples were given via oral gavage. Body weight, food consumption, and water intake were recorded every day. The daily health status of the rats was also monitored. The diarrhea model was performed based on the methods described by van Ampting et al. [12] with slight modifications in antibiotic dosage and days of *Salmonella* infection. As indicated in Figure 1, from day 15 to day 17 of the experiment, rats in the negative control and sample groups were given 100 mg of ampicillin (Sigma-Aldrich, St. Louis, MO, USA; CAS69-52-4) and 15 mg of clindamycin (Goldbio, St. Louis, MO, USA; CAS21462-39-5) which were mixed with basal diet. The morphology of rat feces was recorded, and the fecal DI was determined. From day 18 to day 22 of the experiment, $1 \times 10^9$ CFU of *S. enterica* (strain number: BCRC10747) culture was given to the negative control and sample groups every day. The fecal morphology and diarrhea index were recorded.

**Figure 1.** Schematic diagram of the experimental design in vivo.

### 2.3. Analysis of Fecal Bacteria

After the second day of the experiment, freshly voided feces were collected in an aseptic tube and analyzed by conventional microbiological methods within 2 h. A series of ten-fold dilutions of the homogenized samples were prepared using sterile and anaerobic dilution solution. *B. coagulans*, *Bifidobacterium* spp., *E. coli*, and *Salmonella* were enumerated by spread plating onto different selective mediums, which were glucose yeast extract agar (GYEA), *Bifidobacterium* iodoacetate medium 25 (BIM-25), Levine eosin methylene blue (LEMB) agar (Merck KGaA, Darmstadt, Germany), and mannitol lysine crystal violet brilliant green (MLCB) agar (Cyrusbioscience, New Taipei City, Taiwan), respectively. *B. coagulans* was aerobically cultivated at 37 °C for 48 h [4]. *E. coli* were aerobically cultivated at 37 °C for 72 h and 48 h, respectively. *Salmonella* was aerobically cultivated at 40 °C for 24 h, and *Bifidobacterium* spp. was anaerobically cultivated at 37 °C for 48 h.

The colonies of *B. coagulans* and *Salmonella* isolated in the fecal samples were confirmed genetically by a molecular probing technique using 16S ribosomal RNA gene primers (Supplementary Table S3 and Figures S1 and S2).
2.4. Determination of Fecal Moisture

Fresh rat feces were collected and accurately weighed. The samples were dried in a hot air oven at 80 °C for 6 h and subsequently dried at 105 °C until a constant weight was reached. The fecal moisture content was calculated by the equation:

\[
\text{Fecal moisture content (g/100 g feces)} = \frac{\text{initial weight} - \text{final weight}}{\text{initial weight}}
\]

2.5. Diarrhea Index Assessment

Diarrhea assessments were performed according to the methods described by Sakai et al. [13] with slight modifications. The severity of diarrhea was scored using the following scale, 1: normal (normal and dark brown stool), 2: slight (slightly wet and yellow black stool), 3: moderate (wet and unformed, brown stool), and 4: severe (watery stool, cannot be picked up with tweezers). The incidence of each diarrhea score (1 to 4) and the average diarrhea score were used to evaluate the severity of diarrhea (Figure 2).

![Diarrhea index](image)

**Figure 2.** Morphology of stool in different diarrhea index.

2.6. Determination of SCFAs

Fecal SCFAs were determined according to the method described by Saw et al. [4] with slight modifications. Fresh fecal samples were homogenized with deionized water at a ratio of 1:10 (w/v), followed by centrifugation at 1006 × g for 10 min. Two milliliters of supernatant were combined with 10 μL of isocapric acid (internal standard) and 20 μL of 50% (w/v) sulfuric acid. Diethyl ether was used to extract the SCFAs. The SCFAs in the ether layer (1 μL) were analyzed using Agilent J and WHP-INNOWax GC Column (30 m, 0.25 mm, 0.25 μm). As a carrier gas, helium gas was used with a constant flow rate of 7 mL/min. An Agilent Technologies 7890A system equipped with a flame ionization detector (FID) was used for chromatographic analysis. During the analysis, the initial oven temperature at 80 °C was maintained for 1 min before being raised to 140 °C at a rate of 20 °C/min and held for another 1 min. The temperature was again increased to 220 °C at a rate of 20 °C/min, and lastly, held at 220 °C for 2 min. The temperatures of the detector and the injector were 250 °C and 140 °C, respectively. SCFAs were quantified by comparing the spectrum obtained from the fecal samples of each group with those of standard compounds.
2.7. Histological Assessment of Colonic Inflammation

Rats were sacrificed at the end of the experiment. The procedure described by Aleisa et al. [14] was used to determine the inflammation of colonic tissue. Colon tissue was removed, cut longitudinally, and washed with saline. Fixed colonic tissue (2 cm) in 10% buffered formalin was used for histological evaluation. Embedded specimens were cut into sections of 5 µm thickness in paraffin wax blocks. Colon segments were stained with Hematoxylin and then counterstained with Eosin. Tissue inflammatory infiltration, severity of edema, cellular necrosis, and production of regenerative cells were observed microscopically. Degree of lesions was graded from 1 to 5 depending on severity: 1 = minimal (<1%); 2 = slight (1–25%); 3 = moderate (26–50%); 4 = moderate/severe (51–75%); 5 = severe/high (76–100%).

2.8. Statistical Analysis

Statistical analysis was carried out using the SPSS statistics program (Version 20.0; SPSS, Armonk, NY, USA). All parameters were analyzed by one-way analysis of variance (ANOVA) followed by Duncan’s multiple range test, with the significance level set at \( p < 0.05 \). The data were expressed as mean ± standard deviation (SD).

3. Results and Discussion

3.1. In Vitro Prebiotic Activity

Table 1 shows the prebiotic impact of different carbohydrate components on the growth of \( B. \ coagulans \) BACO-17. The results were compared to the rate of probiotic bacteria growth in the presence of glucose. In this study, among the four carbohydrate samples, the pectin group yielded significantly higher (\( p < 0.05 \)) \( B. \ coagulans \) count than other groups after the sixth hour of cultivation. This indicated that \( B. \ coagulans \) BACO-17 grew faster using pectin as its carbon energy source. A similar growth profile was observed across all prebiotic samples; specifically, the highest bacterial count in the range of 6.83–7.20 Log CFU/mL was observed at the sixteenth hour of incubation. The data suggested that GOS created the best growth condition, followed by FOS.

Table 1. Effects of various carbon sources at the level of 0.5% (w/v) on the growth of \( B. \ coagulans \) (Log cfu/g) in vitro.

| Groups   | 0     | 6     | 9     | 12    | 16    | 24    |
|----------|-------|-------|-------|-------|-------|-------|
| Glucose  | 3.86 ± 0.15 \( ^a \) | 4.69 ± 0.10 \( ^b \) | 5.84 ± 0.20 \( ^a \) | 6.96 ± 0.28 \( ^a \) | 7.23 ± 0.24 \( ^a \) | 7.14 ± 0.14 \( ^a \) |
| XOS     | 3.86 ± 0.15 \( ^a \) | 4.53 ± 0.15 \( ^b \) | 5.36 ± 0.43 \( ^b \) | 6.65 ± 0.67 \( ^a \) | 7.05 ± 0.26 \( ^{ab} \) | 6.96 ± 0.11 \( ^{ab} \) |
| FOS     | 3.86 ± 0.15 \( ^a \) | 4.09 ± 0.17 \( ^d \) | 4.87 ± 0.73 \( ^b \) | 7.03 ± 0.28 \( ^a \) | 7.18 ± 0.12 \( ^a \) | 6.77 ± 0.32 \( ^b \) |
| GOS     | 3.86 ± 0.15 \( ^a \) | 4.29 ± 0.30 \( ^c \) | 5.25 ± 0.80 \( ^b \) | 6.10 ± 0.97 \( ^a \) | 7.20 ± 0.39 \( ^a \) | 7.10 ± 0.30 \( ^a \) |
| Pectin  | 3.86 ± 0.15 \( ^a \) | 4.95 ± 0.14 \( ^a \) | 5.92 ± 0.12 \( ^a \) | 6.45 ± 0.91 \( ^a \) | 6.83 ± 0.16 \( ^b \) | 6.86 ± 0.16 \( ^{ab} \) |

\(^{1} \) Glucose was used as the sole carbohydrate source in the control group. \(^{2} \) In order to select appropriate prebiotics for \( B. \ coagulans \) BACO-17 to create a synbiotic preparation, different prebiotics, including xylo-oligosaccharide (XOS), fructo-oligosaccharide (FOS), galacto-oligosaccharide (GOS), or citrus pectin were used to replace glucose in equal amount. \(^{a-d} \) Values (mean ± SD, \( n = 6 \)) with different superscripts in the same column are significantly different, \( p < 0.05 \).

It was observed that \( B. \ coagulans \) BACO-17 was capable of utilizing all of the tested prebiotics. Noticeably, although \( B. \ coagulans \) BACO-17 initially grew faster using pectin as its energy source, its growth lagged behind other groups after 24 h of cultivation (6.86 Log CFU/mL). This indicated that pectin was not utilized as efficiently as the other prebiotics. The growth rates of \( B. \ coagulans \) BACO-17 in the FOS and GOS groups were similar to that of the glucose control. Our results indicated that GOS might better support the growth of \( B. \ coagulans \) BACO-17. These findings were consistent with the observations made by Cano Roca [15], stating that culturing \( B. \ coagulans \) using GOS led to a more significant change in absorbance value and better coagulation efficiency compared with
that using FOS. Therefore, in this study, FOS and GOS were selected for further evaluation of their prebiotic potential with *B. coagulans* BACO-17 in vivo.

3.2. Food Intake and Body Weight Gain

There were no apparent differences in food intake (24.1–25.8 g/day) and body weight gain (2.3–2.7 g/day) among all animal groups before the intervention (Supplementary Table S1). The daily consumption of the *B. coagulans* BACO-17 and prebiotic samples did not affect healthy rats’ appetite or body weight.

After the antibiotic intervention and *Salmonella* infection, it was found that food intakes in the D (negative control) and samples groups were significantly (*p* < 0.05) reduced. Previous findings from Tulstrup et al. [16,17] have demonstrated that antibiotic therapies cause decreases in body weight and food intake due to antibiotic-perturbed low-diversity microbiota and disturbed appetite. During the diarrhea period, the rate of weight gain in negative control and sample groups was reduced by 16.3–38.7% versus the normal control, with the GB group having the lowest decrease in weight gain (Supplementary Table S1). It has been reported that galacto-oligosaccharide consumption could resist drug-induced body weight loss in a dose-dependent manner [18]. Therefore, it was inferred that combining *B. coagulans* BACO-17 and GOS as a synbiotic could help alleviate antibiotics- and *salmonella*-induced decrease in body weight gain.

3.3. Assessment of Fecal Moisture and Diarrhea Index

Figure 3 illustrates that three days of antibiotic intervention for the animals in the negative control and sample groups starts on day 15. After the antibiotic supplementation, these animals were subsequently infected orally with *S. enterica* culture for 5 days (between day 18 and day 22), followed by another 5 days of the recovery period.

![Figure 3. Change in fecal moisture among different animal groups. Blue zone (days 0 to 14) indicates the period before intervention. Yellow zone (days 15 to 22) indicates the period of antibiotics intervention and *Salmonella* infection. Green zone (days 23 to 25) indicates the recovery period. Mean values on the curves bearing different letters (a–c) indicate significant differences (*p* < 0.05) among different groups on day 25.](image)

Before the antibiotic intervention (from day 0 to day 14), Figure 3 shows no apparent difference in fecal moisture content among all seven dietary groups (57.6–62.2 g/100 g feces). After the supplementation of antibiotics, fecal moisture content of the negative control and five sample groups significantly (*p* < 0.05) increased by about 25% to the highest values (up to 72.5–75.9 g/100 g feces) between day 17 and day 20, followed by a subsequent reduction. During the recovery period (day 25), a significant (*p* < 0.05) reduction in the fecal moisture content was observed only in the BC, GOS, and GB groups (9.0%, 7.3%, and
6.1%, respectively) against their corresponding highest values, implying a better recovery in the animal groups fed either B. coagulans BACO-17 or GOS, alone or in combination.

Regarding the diarrhea index, rats fed a normal diet had diarrhea indexes ranging between 1.13–1.22. After the antibiotic intervention, rats in all dietary groups (except normal control) discharged diarrheic feces, and the severity of diarrhea gradually increased to their highest values (2.66–2.92) between day 17 and day 20 (Figure 4). It appeared that the approach to introducing an antibiotic intervention successfully induced diarrhea symptoms in this study. At the end of this experiment, all five sample groups had a significant ($p < 0.05$) decline in diarrhea indexes, which ranged from 14.5 to 26.9%. Notably, the BC group had the most drastic reduction among these dietary groups (26.9%), followed by the GB group (18.7%). B. coagulans consumption could ameliorate diarrhea symptoms and inhibit gastrointestinal motility in rats [19] and was reported to be able to improve the intestinal environment, defecation frequency, and fecal characteristics in humans [20].

![Figure 4](image_url)

**Figure 4.** Changes in diarrhea index among different animal groups. Blue zone (days 13 to 14) indicates the period before intervention. Yellow zone (days 15 to 22) indicates the period of antibiotics intervention and *Salmonella* infection. Green zone (days 23 to 25) indicates the recovery period. Mean values on the curves bearing different letters (e–h) indicate significant differences ($p < 0.05$) among different groups on day 22. Mean values on the curves bearing different letters (a–d) indicate significant differences ($p < 0.05$) among different groups on day 25.

Interestingly, the BC group had a superior performance in reducing incidences and severity of diarrhea compared with other dietary groups. In sum, using B. coagulans BACO-17 alone and in combination with prebiotics, particularly GOS, could effectively relieve diarrhea.

### 3.4. Changes in Fecal Microflora

In the first two weeks of this feeding experiment (Figure 1), there were no statistical differences in fecal *B. coagulans*, *Bifidobacteria*, and *E. coli* counts among the seven animal groups (Figure 5A–D). These fecal bacterial counts remained stable before the intervention with antibiotics.

Figure 5A reveals that no *Salmonella* was detected in the feces among all different dietary groups in the first two weeks. It indicated that the *Salmonella* count was below the detectable range, although the existence of *Salmonella* was generally expected in the small intestine. The observation was consistent with a previous study claiming that fecal *Salmonella* was undetectable without infection [21]. After the *Salmonella* infection, fecal *Salmonella* counts increased from non-detectable to $3.80–4.03$ Log CFU/g on day 19. However, during the rest of the experiment (from days 19 to 25), no further variation in the fecal *Salmonella* counts ($3.47–4.28$ Log CFU/g) was observed.
Figure 5. Change in different fecal bacteria counts (Log cfu/g) among different animal groups. (A) fecal *Salmonella*; (B) fecal *B. coagulans*; (C) fecal *Bifidobacterium* spp.; (D) fecal *Escherichia coli*. Blue zone (days 0 to 14) indicates the period before intervention. Yellow zone (days 15 to 22) indicates the period of antibiotics intervention and *Salmonella* infection. Green zone (days 23 to 25) indicates the recovery period. Mean values on the curves bearing different letters (e–g) indicate significant differences (*p* < 0.05) among different groups on day 22. Mean values on the curves bearing different letters (a–d) indicate significant differences (*p* < 0.05) among different groups on day 25.

After the antibiotic intervention at day 15, a significant (*p* < 0.05) reduction (ranging from 21.5% to 27.8%) in the fecal *B. coagulans* count was observed in the subsequent 4 days among the BC, FB, and GB groups (Figure 5B), followed by a rebound of bacterial growth on day 19. Although GOS could support a better growth for *B. coagulans* BACO-17 in the in vitro experiment, no apparent difference in the fecal *B. coagulans* count was noted in the in vivo experiment. It was believed that the discrepancy between the in vitro and in vivo experimental results was likely because the in vitro experimental condition was not complex enough to mimic the intestinal microflora in feces and thoroughly examine specific substrates’ prebiotic potential [22].

A similar pattern was also observed in the growth of fecal *Bifidobacteria* in the negative control and sample groups (Figure 5C). The dramatic drop in fecal *Bifidobacteria* counts indicated that antibiotics might affect the growth of fecal bacteria. On day 20 of the experiment, the fecal *Bifidobacteria* counts ascended to a higher level (8.54–9.98 Log CFU/g). It was inferred that continuous *B. coagulans* supplement provided necessary organic acids and other metabolites for the growth of lactic acid bacteria [23] and hence aided in restoring and increasing the total *Bifidobacteria* counts in feces.

From days 15 to 17 of the experiment, it was interesting to note that the intervention with antibiotics resulted in a significant (*p* < 0.05) reduction in the fecal *E. coli* counts in both the negative control group and the other five sample groups (Figure 5D). The phenomenon was probably attributed to the fact that antibiotics suppressed the number of *E. coli* in feces below the detectable range [24]. After the *Salmonella* infection on day 18, the apparently
(p < 0.05) higher fecal E. coli counts versus with their pre-infection state was in agreement with the observations previously reported by Polishchuk et al. [25].

During the recovery period (day 25), Figure 5C,D indicate that the GB group had the highest Bifidobacterium count in feces and the most prominent downward trend in the E. coli counts in feces. It was postulated that the combination of B. coagulans BACO-17 with appropriate prebiotics, particularly GOS, could be explored as a potential symbiotic mixture for improving dysbiosis in feces.

3.5. Changes in Fecal SCFAs

Table 2 presents the concentrations of total SCFAs in rat feces collected on day 14 (before antibiotic intervention), day 19 (after antibiotic intervention and Salmonella infection), and day 25 (end of the experiment). The administration of B. coagulans BACO-17 and the two prebiotics (i.e., FOS and GOS) for 14 days effectively promoted the production of total SCFAs in feces (37.9–64.9%) relative to the normal control. It is worth noting that the supplementation of symbiotic mixtures in GB and FB groups would further result in an approximately two-fold increase (p < 0.05) in the fecal levels of total SCFAs compared with the normal control.

Table 2. The total concentrations of fecal short-chain fatty acids (µmol/g) at different times.

| Groups       | Day 14 (Before Antibiotic Intervention) | Day 19 (After Salmonella Infection) | Day 25 (Recovery Period) |
|--------------|-----------------------------------------|-------------------------------------|--------------------------|
| Normal control | 115.0 ± 20.2 aA | 137.8 ± 23.4 dA | 102.3 ± 23.7 aA |
| Negative control | 124.1 ± 29.3 aC | 52.4 ± 22.1 aA | 94.8 ± 17.6 aB |
| BC | 189.6 ± 27.3 cC | 92.2 ± 6.2 bcA | 145.4 ± 18.1 bb |
| FOS | 170.0 ± 17.9 bcC | 94.9 ± 18.4 cA | 123.3 ± 10.9 abB |
| GOS | 158.6 ± 13.9 bbB | 70.0 ± 25.2 abA | 147.5 ± 38.8 bbB |
| FB | 223.1 ± 11.5 dbC | 67.0 ± 24.5 aA | 145.6 ± 26.8 bbB |
| GB | 232.4 ± 28.0 dcC | 121.5 ± 9.0 dA | 182.1 ± 20.0 dB |

a–d Different lower-case superscripts in the same column indicate significant differences among different groups, p < 0.05. A–C Different capital superscripts in the same row indicate significant differences within the same group, p < 0.05.

After the antibiotic intervention and Salmonella infection, the total concentrations of fecal SCFAs in all animal groups (except normal control) were significantly decreased (p < 0.05). Noticeably, the negative control group had the least total SCFAs (52.4 µmol/g). Mekonnen et al. [26] reported that antibiotic administration could induce an imbalance of gut microbiota and impact carbohydrate metabolism, resulting in lower levels of acetic acid, propionic acid, butyric acid, and total SCFAs in feces. Our findings, as presented in Supplementary Table S2, also demonstrated a dramatic decline in the fecal levels of acetic acid and butyric acid.

During the recovery period (day 25), a general increase in the fecal SCFA concentrations was noted in Table 2. Among the five sample groups, the GB group showed the most remarkable improvement in the elevation of fecal total SCFA levels (182.1 µmol/g), which was almost 2-fold higher than the negative control. More specifically, the BC and GB groups showed a more promising increase in fecal butyric acid concentrations (Supplementary Table S2). Some previous studies have demonstrated that B. coagulans combined with prebiotics remarkably increased intestinal Lactobacillus and Bifidobacterium levels and boosted the production of lactic acid and SCFAs from prebiotics [27,28]. These findings might support the idea of developing a symbiotic by combining B. coagulans BACO-17 and GOS through their enhanced synthesis of fecal SCFAs.

3.6. Histopathological Studies

The H&E staining of colon tissue in each group is shown in Figure 6. The arrowheads indicate submucosae infiltrated with inflammatory cells. The results show that the normal
control group had no apparent inflammatory cell infiltration in the lamina propria. In contrast, the negative control had significantly greater severity of inflammatory cell infiltration (at least a 2-fold increase) and extent of edema than the normal control (Table 3). It is worth noting that the administration of *B. coagulans* BACO-17 alone (BC group) or in combination with GOS (GB, synbiotic group) was more effective in reducing intestinal inflammation (27–31%) and mucosal necrosis (82%) over the negative control. The antibiotics treatment was supposed to cause disturbance of intestinal microbiota and facilitate colonization and infectivity of intestinal pathogens [29].

Figure 6. Cont.
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Table 3. Histopathological scoring of inflammation grading based on H&E staining in different animal groups.

| Groups     | Inflammation  | Edema       | Necrosis     | Regeneration |
|------------|---------------|-------------|--------------|--------------|
| Normal control | 2.00 ± 0.00 a | 0.00 ± 0.00 a | 0.00 ± 0.00 a | 0.00 ± 0.00 a |
| Negative control | 4.33 ± 0.82 c | 1.67 ± 1.21 b | 1.83 ± 0.98 d | 2.00 ± 1.67 bc |
| BC         | 3.17 ± 0.75 b | 1.00 ± 0.89 ab | 0.33 ± 0.52 ab | 0.33 ± 0.52 ab |
| FOS        | 4.33 ± 1.03 c | 1.33 ± 1.03 b | 1.33 ± 1.03 ed | 2.67 ± 2.07 c |
| GOS        | 4.00 ± 0.89 bc | 1.33 ± 1.03 b | 1.00 ± 0.89 bcd | 2.33 ± 1.86 c |
| FB         | 3.67 ± 0.52 bc | 0.67 ± 0.52 ab | 0.67 ± 0.52 abc | 1.33 ± 1.37 abc |
| GB         | 3.00 ± 1.10 bc | 0.67 ± 0.82 ab | 0.33 ± 0.82 ab | 0.33 ± 0.82 ab |

ab Values (mean ± SD, n = 6) with different superscripts in the same column are significantly different, p < 0.05.

This indicated that antibiotic administration and Salmonella infection have successfully caused colon injury and cell damage in rats. As presented in Figure 5A,D, it is believed that the inflammatory symptoms caused by Salmonella infection not only promoted the growth of Salmonella in feces but also facilitated the proliferation of fecal commensal bacteria (such as E. coli) [30]. Compared with the negative control group, GB and BC groups presented less inflammation infiltration in mucosal lamina propria and submucosa and less cell necrosis in colonic tissues (Figure 6). It was postulated that the apparent improvement observed in BC and GB groups might be partly attributed to the remarkable elevation in the fecal butyric acid concentrations. Alonso and Guarner [31] suggested that butyrate might serve as a primary energy source for colon cells, achieve anti-inflammatory effects by inhibiting the NF-κB pathway and reducing the expression of pro-inflammatory genes, enhance the integrity of the intestinal mucosal barrier, and regulate apoptosis.

The short-chain fatty acids produced during digestion might help maintain the intestinal mucosal barrier and exert immune regulation. Specifically, high acetic acid and butyric acid concentrations could exert potent anti-inflammatory effects [32]. It was speculated that the relatively higher fecal concentrations of butyric acid in BC and GB groups might contribute to reducing the extent and severity of colon inflammation in these groups. These results suggested that B. coagulans BACO-17 alone or in combination with GOS could effectively relieve intestinal inflammation.

4. Conclusions

Intestinal dysbiosis triggered by the antibiotic treatment could result in diarrhea. In the BC and GB groups, it was found that the beneficial bacteria counts and fecal SCFA concentrations were increased; meanwhile, the diarrhea symptoms were ameliorated. However, none of the sample groups effectively inhibited Salmonella colonization in the intestine.
It was speculated that the antibiotics intervention had disturbed gut microbiota balance and caused inflammation and that *Salmonella* had competitive advantages over the commensal microbiota under such a condition. Based on the results of this study, the preventive effects of *B. coagulans* BACO-17 alone or combined with GOS as a synergetic mixture on intestinal inflammatory diseases could be a promising adjuvant therapy; however, clinical trials are needed to further validate the therapeutic effects. Further studies are also needed for a more complete understanding of the changes in microbial biofilm associated with intestinal tissue and complex interactions between the host and gut microbiota.

**Supplementary Materials:** The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/pr10102123/s1, Figure S1: Agarose gel electrophoresis of the amplified target of 16S rRNA gene. (A) *Bacillus coagulans* BACO-17, (B) *Salmonella enterica* subsp. *enterica* BCRC 10747; Figure S2: Partial sequence of 16S rRNA gene and top listed BLAST results. (A) *Bacillus coagulans* BACO-17, (B) *Salmonella enterica* subsp. *enterica* BCRC 10747; Table S1: Comparisons of the average food intake, water intake, and body weight gains before and after intervention among different animal groups; Table S2. (A–C) The concentrations of acetic acid (µmol/g) in fecal samples collected at different time points; Table S3: Primers used for 16S rRNA gene analysis [33,34].

**Author Contributions:** Conceptualization, C.-F.C., M.-Z.W., Y.-W.H. and W.-J.C.; methodology, M.-Z.W., Y.-W.H., T.-C.S., H.-F.C. and C.-F.C.; formal analysis, M.-Z.W., Y.-W.H., T.-C.S. and C.-Y.L.; data curation, M.-Z.W., Y.-W.H., T.-C.S., H.-F.C., C.-Y.L. and C.-F.C.; writing-original draft preparation, M.-Z.W., Y.-W.H., Y.-C.W., T.-C.S. and C.-F.C.; writing-review and editing, C.-F.C., M.-Z.W., Y.-C.W. and C.-Y.L.; supervision, C.-F.C.; project administration, C.-F.C.; funding acquisition, C.-F.C. and W.-J.C. All authors have read and agreed to the published version of the manuscript.

**Funding:** This research was supported by the National Science and Technology Council of the Republic of China (MOST 111-2320-B-005-004-MY2).

**Data Availability Statement:** Data is contained within the article.

**Conflicts of Interest:** The authors declare no conflict of interest.

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