<table>
<thead>
<tr>
<th>Title</th>
<th>Ameliorating Effects of Short-Chain Inulin-Like Fructans on the Healing Stage of Trinitrobenzene Sulfonic Acid-Induced Colitis in Rats</th>
</tr>
</thead>
<tbody>
<tr>
<td>Author(s)</td>
<td>Hino, Shingo; Ito, Hiroyuki; Bito, Hiroyuki; Kawagishi, Hirokazu; Morita, Tatsuya</td>
</tr>
<tr>
<td>Citation</td>
<td>Bioscience, Biotechnology, and Biochemistry. 75(11), p. 2169-2174</td>
</tr>
<tr>
<td>Issue Date</td>
<td>2011-11-07</td>
</tr>
<tr>
<td>URL</td>
<td><a href="http://hdl.handle.net/10297/6928">http://hdl.handle.net/10297/6928</a></td>
</tr>
<tr>
<td>Rights</td>
<td>Copyright © 2011 by Japan Society for Bioscience, Biotechnology, and Agrochemistry</td>
</tr>
</tbody>
</table>
Ameliorating Effects of Short-Chain Inulin-Like Fructans on the Healing Stage of Trinitrobenzene Sulfonic Acid-Induced Colitis in Rats

Shingo Hino, Hiroyuki Ito, Hiroyuki Bito, Hirokazu Kawagishi, and Tatsuya Morita

Department of Applied Biological Chemistry, Faculty of Agriculture, Shizuoka University, 836 Ohyaa, Shizuoka 422-8529, Japan

Graduate School of Science and Technology, Shizuoka University, 836 Ohyaa, Shizuoka 422-8529, Japan

Received June 15, 2011; Accepted August 4, 2011; Online Publication, November 7, 2011

We evaluated the ameliorating effects of short-chain inulin-like fructans (SIF) with different degrees of polymerization (DP) on the healing stage of trinitrobenzene sulfonic acid (TNBS)-induced colitis in rats. The rats were assigned to 3 groups 10 d after the colitis induction, and fed for 24 d on a control diet or diet including 60 g of DP4 or DP8/kg. The fecal myeloperoxidase (MPO) activity and IgA were measured at the end of the study. DP4, but not DP8, significantly reduced the colonic inflammation accompanied by higher cecal concentrations of short-chain fatty acids, propionate in particular, and lactic acid-producing bacteria. DP4 therefore accelerated the healing process of TNBS-induced colitis, even when the treatment was initiated after inducing colitis.

Key words: short-chain inulin-like fructans; short-chain fatty acids; lactobacilli; trinitrobenzene sulfonic acid (TNBS) colitis; rats

Inflammatory bowel disease is a chronic disease of the digestive tract and usually refers to two related conditions, ulcerative colitis and Crohn’s disease, characterized by spontaneously relapsing inflammation. The etiology of inflammatory bowel disease remains unclear, but it is believed that genetic, immunologic and colonic environmental factors, including microbiota, may be involved. Diet is a major factor affecting the luminal environment and intestinal immune function. Considerable attention has recently been paid to prebiotic and probiotic treatments which aim to stimulate the growth of lactic acid-producing bacteria, reduce colonic pH, and maintain mucosal integrity by augmenting the production of short-chain fatty acids (SCFAs). Prebiotic and probiotic treatments also have the potential to increase both luminal mucin and IgA that are related to the mucosal barrier function against bacteria and endotoxin translocation.

We have previously shown that pre-treating short-chain inulin-like fructans (SIF), with average degrees of polymerization (DP) of 4 (virtually the same as fructooligosaccharides) and 8, exerted preventive effects on the acute phase of trinitrobenzene sulfonic acid (TNBS)-induced colitis in rats. SIF, and DP8 in particular, increased the cecal concentrations of SCFA, mucin and IgA before the induction of colitis, and reduced bacterial and endotoxin translocation to the mesenteric lymph node and portal blood 3 d after the TNBS administration. However, these beneficial effects of DP4 and DP8 on the cecal valuables were abolished 10 d after the TNBS administration, despite the significant reduction of colonic inflammation. We therefore assumed that these anti-inflammatory effects of SIF might be exerted through a shield-like effect against endotoxin and bacterial translocation at the early stage of colitis, leading to the reduced colonic damage 10 d after the induction of colitis.

A number of studies examining the effects of prebiotics, including dietary fiber and probiotics, on TNBS-induced rat colitis have already been reported, but most of them evaluated the preventive effects in the acute phase of colitis within 1–2 weeks after TNBS administration; the outcome was mostly successful when the treatment began before the induction of colitis. Indeed, Peran et al. have reported for rats that a pre-treatment of Lactobacillus rutei and L. fermentum, starting 2 weeks before TNBS administration, significantly reduced colonic damage both histologically and biochemically. However, Kennedy et al. have shown that probiotic Lactobacillus plantarum species 299 and oat fiber had no effect on colonic inflammation when given for 7 d following TNBS administration. This was probably because the inflammation in the acute phase of TNBS colitis was so extensive and severe due to the caustic properties of TNBS and ethanol, besides the hapten-induced immunological modulation. Even chemotherapy, using an inhibitor of neutrophil activation and dexamethasone, did not produce any significant effects when the treatment was initiated after the induction of colitis. Taken together, it is likely that, apart from the preventive effect, the acute phase of TNBS colitis is not suitable for examining any beneficial (therapeutic) effects of dietary factors.

It has been well established that the rat TNBS colitis model induced long-lasting colitis for at least 6 weeks after its induction, although the colonic inflammation gradually healed and damage scores decreased from its...
inflammatory peak 7 d after the induction.1,4,18) This may permit us to study the influence of dietary factors during the healing stage of colitis. The present study was accordingly aimed to examine whether the effects of SIF could accelerate the healing process of TNBS colitis in rats. Once colitis had been established, the rats were assigned to each dietary group on the basis of the severity of colonic inflammation as assessed by the fecal myeloperoxidase activity (MPO). The dietary treatment was started 10 d after the induction of colitis and continued for 24 d.

Materials and Methods

Materials. SIF with DP4 and DP8 were used. Metolig® P DP4 was purchased from Meiji Seika (Tokyo, Japan), its composition being 44% of 1-kestose, 46% of nystose, and 10% of 1-f-fructofuranosyl nystose. DP8 (with a 5–13 range of DP) was prepared by enzymatic synthesis, using a novel fungal enzyme β-fructofuranosidase from bacillus sp. 217C-11, as described previously.3,19)

Care of the animals. Male Sprague-Dawley rats (SPF) were purchased from Shizuoka Laboratory Animal Center (Hamamatsu, Japan). They were housed in individual screen-bottomed stainless steel cages in a room with controlled temperature (23 ± 2 °C) and lighting (on from 8:00 to 20:00). The rats were fed with a control diet for at least 3 d for adaptation which was formulated from 250 g/kg of casein, 652.25 g/kg of cornstarch, and 50 g/kg of corn oil.19) The rats were administered with a control diet for 7 d, the rats were lightly anesthetized with diethyl ether and then treated with an intracolic injection of a 20% glycerin solution (0.2 mL/rat), using a lubricated polypropylene catheter (1.5 mm diameter) inserted 8 cm into the colon via the anus. Preliminary results indicated that this treatment was useful for removing the colonic contents. Defecation was usually completed within 15 min, and the colon was kept empty for at least 60 min after the treatment. At 60 min after administering the glycerin solution, 8 rats were anesthetized with diethyl ether and then treated with an intracolic instillation and measured for the MPO activity. The rats were then anesthetized with diethyl ether 8 d after the instillation, and the colon and blood were collected. The colon was cut open longitudinally and the colonic contents removed. The colonic tissue was stored at −80 °C pending measurement of the MPO activity. The blood sample was obtained from the abdominal aorta with a heparinized syringe. The plasma was then obtained, after centrifugation at 2000 × g for 10 min at 4 °C, and used for measuring α1-AG.

Ameliorating effects on the healing stage of TNBS-induced colitis by SIF ingestion. Thirty-five rats (207–237 g, 7 weeks old) were used. After feeding the control diet for 7 d, the rats were administered with 20% glycerin and then TNBS/50% ethanol in the same manner as that used in preliminary experiment 1. The feces were collected 6–9 d after instillation and measured for MPO activity (6–7 d) and IgA concentration. The rats were then anesthetized with diethyl ether at the end of test period, and the cecum and the colon were excised. The cecal contents were removed, weighed, and divided into two portions. One was freeze-dried and used for a mucin analysis, and the other was used for measuring the pH value, organic acids, IgA, bifidobacteria and lactobacilli. The colon was handled in the same manner as that described for the preliminary experiment.

MPO activity. The colonic MPO activity was determined as previously described.25) Briefly, the colon was minced and homogenized by a Polytron device in a 50 mm potassium phosphate buffer (pH 6.0) containing 0.5% hexadecltrimethyl ammonium bromide. The resulting homogenate was subjected to three cycles of freeze-thawing and sonication, and then centrifuged at 20,000 × g for 30 min. The feces were handled in the same manner as that described for the colonic tissue, but without freeze-thawing and sonication. The supernatants of the colonic and fecal homogenates were used to determine the MPO activity, using 0.0005% hydrogen peroxide as the substrate for MPO. One unit of MPO activity is defined as that converting 1 μmol of hydrogen peroxide to water per 1 min at 25 °C.

Plasma α1-AG. Plasma α1-AG was analyzed by an α1-acid glycoprotein rat ELISA kit (Life Diagnostics, PA, USA) according to manufacturer’s instructions.

Cecal pH and organic acids. After the cecal contents had been homogenized, a portion of the homogenate was diluted with the same weight of distilled water, and then the cecal pH value was measured with a C-1 compact pH meter (Horiba, Tokyo, Japan). The cecal organic acids were measured by the internal standard method25) with an LC-10A HPLC instrument (Shimadzu, Kyoto, Japan) equipped with a Shim-pack SCR-102H column (8 mm i.d. × 30 cm long, Shimadzu) and a CDD-6A electro-conductivity detector (Shimadzu).

Secretory IgA. The feces and cecal contents were homogenized with a 10-fold volume of phosphate-buffered saline (PBS, pH 7.4) containing Complete protease inhibitors (Roche, Basel, Switzerland) for feces and 0.02 M sodium azide and 0.5% Tween20 for the cecal contents. Each of these homogenates was centrifuged at 15,000 × g, and the resulting supernatant was applied to an IgA determination by ELISA, using Nunc-immuno MaxiSorb F96 plates with a small modification25) of the method described by Grewal et al.25) Each assay was conducted in duplicate.

O-Linked oligosaccharide chains. Mucins were extracted by the method of Bovee-Oudenhoven et al.24) with some modifications.19) The O-linked oligosaccharide chains were determined as a mucin marker after appropriately diluting the mucin fraction, the O-linked oligosaccharide chains were measured with a fluorimetric assay26) that discriminated O-linked glycoproteins (mucin) from N-linked glycoproteins, as described by Bovee-Oudenhoven et al.24) Standard solutions of N-acetyl galactosamine (Sigma, St. Louis, MO, USA) were used to calculate the quantity of oligosaccharide chains liberated from mucins during the procedure.

Isolation of bacterial DNA. A 50 mg amount of the cecal contents was suspended in 1 mL of sterile PBS (pH 7.2), and the suspension centrifuged at 700 × g for 1 min. The supernatant was collected and re-centrifuged at 10,000 × g for 10 min to yield the bacterial fraction. Bacterial DNA was isolated from this fraction by Isolplant (Nippon Gene, Tokyo, Japan) according to manufacturer’s instructions. The concentration and purity of the bacterial DNA were estimated by measuring its OD260/280 value.

Real-time PCR for bifidobacteria and lactobacilli. Amplification and detection of the bacterial DNA were performed with an ST-300 Light Cycler (Roche). Bifidobacterium genus-specific (forward, TCG CCT C/TC TG T GA AAG; reverse, CCA CAT CCA CCA/GA/TCC AC)26) and Lactobacillus genus-specific (forward, TGG AAA CAG (A/G) TG CTA ATG CCG; reverse, GTC CAT TGT GGA AGA TTC CC)25) primer pairs were used. Real-time PCR was performed in a reaction volume of 20 μL, the mixture containing 10 μL of SYBR
Prenix Ex Taq II (Takara Bio, Tokyo, Japan), 0.8 μL of 200 nM each of the forward and reverse primers, 6.4 μL of H₂O and 2 μL of the bacterial DNA templates. The reaction conditions were 95 °C for 30 s, then 43 cycles at 95 °C for 5 s and 64 °C for 30 s for quantifying the bifidobacteria, and 95 °C for 30 s, then 43 cycles at 95 °C for 5 s and 65 °C for 30 s for quantifying the lactobacilli. The amplification specificity was confirmed by subjecting the PCR products to a melting curve analysis. Bifidobacterium animals (JCM 1190) and Lactobacillus murinus (JCM 171) were cultured in De Man, Rogosa and Sharpe broth (Becton Dickinson, Rockville, MD, USA), and genomic DNA was extracted by Isolplant-II (Nippon Gene) according to the manufacturer’s instructions. Fragments of 16S rDNA were amplified with a QIAprep Spin Miniprep kit (Qiagen, Germantown, MD, USA) and used as a standard for real-time PCR. Escherichia coli and Bifidobacterium were picked out and grown in Luria–Bertani broth. Plasmid DNA was extracted with a GFXe PCR DNA and Gel Band Purification kit (GE Healthcare). Bifidobacterium and Lactobacillus were only apparent between the DP8 and control diet groups (control, DP4 and DP8 diets than by those fed the control diet). The respective food intake and body weight gain were 156 ± 5 g/day and 40 ± 3 g/day for the normal rats, and 107 ± 4 g/day and 11 ± 3 g/day for those on the TNBS-administered diet. The colonic MPO activities were about 6-fold higher in the TNBS-administered rats than in the normal rats (normal, 4.2 ± 0.8; TNBS, 25.2 ± 3.5 unit/g of tissue). Both the fecal MPO activity (normal, 0.05 ± 0.01; TNBS, 0.09 ± 0.01 unit/g of feces) and the plasma α1-AG concentration (normal, 0.14 ± 0.00; TNBS, 0.20 ± 0.01 mg/mL) were also significantly higher in the TNBS-administered rats than in the normal rats. The correlations between the colonic MPO activity and both the fecal MPO activity and plasma α1-AG concentration were significant (Fig. 1a and b). However, higher correlation was apparent between the fecal MPO activity and colonic MPO activity. Even if the data from normal rats are omitted, significant correlation was apparent between the fecal MPO activity and colonic MPO activity with a correlation coefficient of 0.81.

**Results**

**Fecal MPO and plasma α1-AG as non-invasive inflammation markers (preliminary experiment)**

The average body weight of the rats just before the TNBS administration was 278 ± 2 g. The food intake and body weight decreased sharply for the first few days after the TNBS administration and then gradually recovered in the following days. The average body weight of the rats was 286 ± 3 g before the dietary treatment. The food intake was significantly lower during the dietary treatment by the rats fed with the DP4 and DP8 diets than by those fed the control diet (control, 487 ± 9; DP4, 444 ± 9; DP8, 387 ± 13 g/24 d), although significant differences in the body weight gain were only apparent between the DP8 and control diet groups (control, 113 ± 2; DP4, 103 ± 5; DP8, 75 ± 7 g/24 d; Fig. 2a and b). Soft unformed feces were observed during the initial few days from the rats fed with the DP4 and DP8 diets.

At the start of the dietary treatment (d 0), the average fecal MPO activity among the rats was 0.9 ± 0.3 unit/g of feces. The fecal MPO activity in the rats fed the DP4 diet was increased by d 7, then decreased to the same level as that in the control group by d 14, and remained constant until the end of the test period (Fig. 3a). The fecal MPO activity in the rats fed the DP8 diet was significantly higher than in those fed the control diet by d 7 and d 14, and was comparable with those on the control diet by d 21. The fecal IgA concentration was significantly higher in the rats fed the DP4 and DP8 diets than in those fed the control diet by d 7 (Fig. 3b).

The autopsy showed no ulcer and bleeding when macroscopically observed in the colonic mucosa of all

**Fig. 1.** Correlation of the Colonic MPO Activity with the Fecal MPO Activity (a) and the Plasma α1-Acid Glycoprotein Concentration (b) in Rats by 8 d after TNBS Administration.

Solid lines incorporate all of the data. Broken lines omit data from the normal rats. ●, TNBS-treated rats; ○, normal rats.
of the dietary groups. However, the relative colon weight differed among the groups, being highest in the DP8 group, intermediate in the control group, and lowest in the DP4 group (Table 1). The colonic MPO activity was significantly lower in the DP4 group than in the control and DP8 groups. The cecal tissue weight was highest in the DP8 group, intermediate in the DP4 group and lowest in the control diet group. The weight of the cecal contents was significantly higher in the rats fed the DP4 and DP8 diets than in those fed the control diet, and a significant decline of the cecal pH value was apparent in the rats fed the DP4 and DP8 diets. The cecal concentrations of organic acids differed significantly among the groups, the concentrations of acetate, propionate and total SCFA being highest in the rats fed the DP4 diet. The concentrations of n-butyrate and succinate were higher in the rats fed the DP4 and DP8 diets than in those fed the control diet, and the lactate concentration was highest in the rats fed the DP4 and DP8 diets. The cecal bifidobacteria were detected in some of the rats fed the DP4 (6/10) and DP8 (5/10) diets, but not detected in the rats fed the control diet (0/15).

Discussion

It is a prerequisite that the average severity of colonic inflammation among the dietary groups is comparable at the beginning of the dietary treatment when evaluating the effects of dietary factors on the healing stage of colitis. Saiki and Wagner et al. have shown in a human experiment that the stool level of MPO in patients with active inflammatory bowel disease was correlated with laboratory parameters and the endoscopic grade of inflammation. The present results also showed that the fecal MPO activity was correlated with that of the inflamed colonic tissues in rats with TNBS colitis. Accordingly, after colitis had been established, the rats were assigned to each dietary group on the basis of the severity of the fecal MPO activity.

Unlike the results from a previous study showing that both DP4 and DP8 had preventive effects against the development of TNBS-induced colitis in rats, the present results clearly show that only DP4, but not DP8, ameliorated the healing process of TNBS-induced colitis when the dietary treatment was initiated 10d after the induction of colitis. This ameliorating effect of DP4 was accompanied by higher cecal concentrations of SCFA, propionate in particular, and by the cecal numbers of lactobacilli and bifidobacteria when compared with...
those of the control and DP-8 diets. The present study is, to our knowledge, the first to demonstrate an ameliorating effect of fructooligosaccharides (DP4) on colonic inflammation when the treatment was initiated after establishing colitis.

SCFAs, the predominant end-products of non-digestible carbohydrate fermentation in the colon, exert a variety of physiological effects. Inter alia, butyrate is the major energy source of colonocytes and inhibits the activation of nuclear factor-{kappa}B that encodes proteins playing important roles in immune response and inflammation. The physiological concentration of butyrate has also caused a significant increase in mucus secretion in the colonic lumen that may serve as an anti-inflammatory agent. Butyrate has also caused a significant increase in mucus secretion in the colonic lumen that may serve as an anti-inflammatory agent. Butyrate enemas have been shown to reduce, but rather aggravated colonic inflammation in the previous study, 7 the DP8 treatment failed to decrease colonic hypersensitivity and inflammation. Despite the higher butyrate and propionate concentrations of cecal propionate and acetate in the rats fed both the DP4 and DP8 diets, no effect on colonic inflammation was apparent in rats fed the DP8 diet. We therefore cannot ascribe the healing effect of DP4 solely to the higher butyrate concentration.

SCFAs, and particularly acetate and propionate, have recently been found to bind and activate G-protein-couple receptor 43 (GPR43, also known as FFAR2) present mainly in neutrophils. It has also been suggested that the stimulation of GPR43 by SCFAs could play a central role in the normal resolution of colonic inflammatory responses. In this respect, it is notable that the cecal concentrations of acetate and propionate were much greater in rats fed the DP4 diet than in those fed the control and DP8 diets. It is therefore plausible to assume that, under the experimen
tal conditions used in the present study, the higher concentrations of cecal propionate and acetate in the rats fed the DP4 diet might have been responsible for the reduction of colonic inflammation during the healing process of TNBS colitis.

Another possibility for the anti-inflammatory effects of DP4 may have been the increased numbers of cecal lactobacilli and bifidobacteria. Cherbut et al. have suggested that the capacity of fructooligosaccharides (virtually DP4) to increase the lactobacilli count was the main mechanism explaining their protective effect on colonic inflammation, rather than the increased SCFA production. Indeed, lactic acid bacteria have evoked a local immune stimulus to increase the levels of luminal secretory IgA and anti-inflammatory cytokine-like IL-10. Ingesting DP4 increased the cecal concentration of lactic acid-producing bacteria accompanying the increased cecal IgA concentration to the greater degree than that observed in the control and DP8 groups.

Despite the greater preventive effects shown by DP8 in the previous study, the DP8 treatment failed to reduce, but rather aggravated colonic inflammation in the healing stage of colitis. The reason for this has not yet been fully elucidated, although it is well known that administering TNBS altered the microflora, decreasing lactobacilli and increasing gram-negative bacilli in the rat cecum. The bacterial capacity to ferment ingested SIF under such conditions might be considerably compromised when compared with normal conditions just before the induction of colitis. Our previous study has clearly shown that the beneficial effects of DP4 and DP8 on cecal SCFAs, lactate, pH value, mucin and IgA were totally abolished by 10 d post-TNBS administration, even when the treatment started before the induction of colitis. However, the importance of the substrate size (DP) in the fermentation of fructans has been reported in a previous study, indicating that 1-kestose and nystose were directly incorporated into

---

**Table 1.** Colonic and Cecal Variables in Rats with TNBS-Induced Colitis after Ingestion of the Respective Diets for 24 d

<table>
<thead>
<tr>
<th>Variable</th>
<th>Control</th>
<th>DP4</th>
<th>DP8</th>
</tr>
</thead>
<tbody>
<tr>
<td>Colonic Length, cm</td>
<td>11.3 ± 1.3</td>
<td>11.7 ± 0.4</td>
<td>12.1 ± 0.4</td>
</tr>
<tr>
<td>Relative weight, mg/cm²</td>
<td>120 ± 7³⁰</td>
<td>108 ± 6³⁰</td>
<td>148 ± 13³⁰</td>
</tr>
<tr>
<td>MPO activities, unit/g of colon</td>
<td>8.1 ± 1.4³⁰</td>
<td>3.7 ± 0.9³⁰</td>
<td>7.4 ± 1.5³⁰</td>
</tr>
<tr>
<td>Cecum</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Tissue, g</td>
<td>0.7 ± 0.0³⁰</td>
<td>1.2 ± 0.1³⁰</td>
<td>2.6 ± 0.2³⁰</td>
</tr>
<tr>
<td>Contents¹, g</td>
<td>2.0 (1.3–2.8)</td>
<td>3.7 (2.6–5.5)</td>
<td>3.1 (0.5–4.3)</td>
</tr>
<tr>
<td>pH</td>
<td>7.6 ± 0.0³⁰</td>
<td>6.8 ± 0.1³⁰</td>
<td>6.4 ± 0.1³⁰</td>
</tr>
<tr>
<td>Organic acids, µmol/g of wet contents</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Acetate¹</td>
<td>47 (23–55)</td>
<td>75 (42–102)</td>
<td>54 (20–132)</td>
</tr>
<tr>
<td>Propionate¹</td>
<td>12 (6–16)</td>
<td>46 (10–61)</td>
<td>9 (4–55)</td>
</tr>
<tr>
<td>-Butyrate</td>
<td>7 ± 3³⁰</td>
<td>25 ± 3³⁰</td>
<td>20 ± 4³⁰</td>
</tr>
<tr>
<td>Total SCFAs²⁻²³</td>
<td>61 (30–118)</td>
<td>145 (64–197)</td>
<td>79 (30–232)</td>
</tr>
<tr>
<td>Lactate¹</td>
<td>0 (0–1)</td>
<td>0 (0–21)</td>
<td>9 (0–40)</td>
</tr>
<tr>
<td>Succinate¹</td>
<td>2 (1–6)</td>
<td>18 (9–35)</td>
<td>5 (1–19)</td>
</tr>
<tr>
<td>IgA, µg/g of wet contents</td>
<td>46 ± 8³⁰</td>
<td>229 ± 60³⁰</td>
<td>129 ± 34³⁰</td>
</tr>
<tr>
<td>Mucin¹, µmol/g of wet contents</td>
<td>0.3 (0.2–0.7)</td>
<td>0.5 (0.4–0.7)</td>
<td>1.3 (0.5–2.6)</td>
</tr>
<tr>
<td>Lactobacilli¹, ×10⁸ copies/g of wet contents</td>
<td>0.9 (0.2–2.6)</td>
<td>3.5 (0.2–123)</td>
<td>1.7 (0.2–3.4)</td>
</tr>
<tr>
<td>Bifidobacteria¹, ×10¹⁰ copies/g of wet contents</td>
<td></td>
<td>23.1 (6.8–619.7)</td>
<td>0.5 (1.9–2.8)</td>
</tr>
</tbody>
</table>

Data are expressed as the mean ± SE or median (range); control, n = 15; DP4, n = 10; DP8, n = 10. Values not sharing a common superscript letter are significantly different when analyzed by the Tukey-Kramer test.

³ Data were examined by Kruskal-Wallis one-way ANOVA, followed by the Kolmogorov-Smirnov two-sample test. *p < 0.05 vs. control.

¹ Sum of acetate, propionate, and -butyrate.

² Detected number of rats; p < 0.05 vs. DP8 by the Mann-Whitney U-test.

³ Not detected.
Lactobacillus paracasei without any hydrolysis and were rapidly fermented.\textsuperscript{44} Since DP4 mainly consisted of 1-kestose and nystose, these fractions might be preferable as fermentation substrates by lactobacilli, rather than DP8 with higher DP. Differences in the fermentability by lactobacilli between DP4 and DP8 in the damaged condition might therefore explain the different concentrations of cecal acetate and propionate, and the numbers of lactic acid-producing bacteria after the prolonged treatment of these SIFs for 24 d, resulting in the different outcomes of their anti-inflammatory effects.

Both DP4 and DP8 induced a transient increase in fecal MPO activities during the initial stage of 7–14 d after the dietary treatment, although the reason for this remains unclear. However, TNBS administration per se has promoted the proliferation of gram-negative bacilli during the acute phase of colitis.\textsuperscript{42,43} It is therefore reasonable to assume that expansion of the pool size of the cecal endotoxin may result in transient exacerbation of the colonic inflammation.\textsuperscript{43} However, DP4 is the most effective as a probiotic strain compared to DP8 with higher DP load from gram-negative bacilli by the shorter period of the treatment was started after the induction of colitis.\textsuperscript{42,43} It is therefore reasonable to assume that expansion of the pool size of the cecal endotoxin may result in transient exacerbation of the colonic inflammation.\textsuperscript{43} However, DP4 is the most effective as a probiotic strain compared to DP8 with higher DP.

In conclusion, DP4 alone, but not DP8 ameliorated the healing process of TNBS-induced colitis, even when the treatment was started after the induction of colitis. The higher concentration of cecal propionate and the stimulated proliferation of lactic acid-producing bacteria resulting from DP4 ingestion might be closely associated with the ameliorating effect on the healing process for TNBS colitis in rats.

References