Probiotics Isolated From Animals in Northwest China Improve the Intestinal Performance of Mice

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Antibiotic resistance is an increasingly prevalent problem worldwide. Probiotics are live microorganisms that provide health benefits to human beings and animals and also antimicrobial activity against pathogens and might be an antibiotic alternative. The gastrointestinal tract of animals can be a suitable source of finding novel antimicrobial agents, where the vast majority of gut microbes inhabit and a plurality of antimicrobial producers exhibit either a wide or narrow spectrum. Animals that live in Northwest China might possess a special commensal community in the gut. Therefore, the purpose of this study was to assess the effects of three probiotic strains (including *Lactobacillus salivarius* ZLP-4b from swine, *Lactobacillus plantarum* FBL-3a from beef cattle, and *Bacillus velezensis* JT3-1 from yak), which were isolated from livestock in this area, on the overall growth performance, immune function, and gut microbiota of mice. The results showed that the *L. salivarius* ZLP-4b group not only improved the growth performance but also amended the intestinal mucosa morphology of mice. Furthermore, the supplementation of *L. plantarum* FBL-3a and *L. salivarius* ZLP-4b strains significantly increased the content of anti-inflammatory cytokines IL-4 and IL-10 but decreased the pro-inflammatory factor IL-17A. The levels of pro-inflammatory factors IL-6, IL-17A, and TNF-α were also decreased by the *B. velezensis* JT3-1 group pretreatment. The 16S rDNA sequence results showed that the probiotic administration could increase the proportion of Firmicutes/Bacteroidetes intestinal microbes in mice. Furthermore, the relative abundance of *Lactobacillus* was boosted in the JT3-1- and ZLP-4b-treated groups, and that of opportunistic pathogens (including Proteobacteria and Spirochaetes) was diminished in all treated groups compared with the control group. In conclusion, *B. velezensis* JT3-1 and *L. salivarius* ZLP-4b supplementation enhanced the overall performance, intestinal epithelial mucosal integrity, and immune-related cytokines and regulated the intestinal microbiota in mice.

Keywords: *Bacillus velezensis*, *Lactobacillus plantarum*, *Lactobacillus salivarius*, immunity, intestinal flora
INTRODUCTION

Antibiotic has been applied for almost 100 years as a predominant strategy in controlling infectious diseases and improving the growth performance of animals (1). Along with the inappropriate and excessive use of antibiotics, antibiotic resistance issues have been gradually exposed (2). There is an increasing number of bacteria which have developed drug resistance, such as Salmonella with multidrug resistance (3). Besides this, antibiotic resistance is also found in association with gut microbiota disturbance. Furthermore, dysbiosis of microbiota might cause numerous diseases, including obesity, autoimmune diseases, allergy, and intestinal diseases such as irritable bowel syndrome (IBD) (4). Consequently, alternatives to antibiotic treatment were required to be explored (5).

Probiotics have been proven to confer benefits to human and animals within reasonable adoption (6). At present, major probiotics can be classified into Lactobacillus spp. (such as Lactobacillus rhamnosus and Lactobacillus acidophilus), Bacillus spp. (Bacillus subtilis), Bifidobacterium spp. (Bifidobacterium longum and Bifidobacterium animalis), yeast (Saccharomyces cerevisiae) and Clostridium spp. (Clostridium butyricum), and so on (7). In addition, next-generation probiotics (Akkermansia muciniphila, Faecalibacterium prausnitzii, and Eubacterium hallii) and genetically modified probiotics (GM probiotics: mutation and overexpression) were vigorously researched (8, 9). The basic mechanism by which probiotics exert beneficial effects is explained as follows: (1) colonization and restoration of disordered intestinal microbiota in the host, (2) competitive exclusion of harmful microbes and antimicrobial molecule production, (3) cell antagonism, cell adhesion, and mucin expression, and (4) regulation of innate and acquired immunity of the host (10). Studies demonstrated that B. subtilis supplementation could ameliorate heat-induced behavioral and inflammatory reactions (11). Sun et al. (12) reported that Bifidobacterium administration altered the commensal community in the gut of mice and regulated the mucosal immunity as determined by Tregs in the colitis model under challenge (13). Therefore, this study set out to assess the effects of Bacillus spp. (Bacillus velezensis JT3-1 obtained from yak) and the effects of two strains of Lactobacillus (Lactobacillus plantarum FBL-3a sourced from beef cattle and Lactobacillus salivarius ZLP-4b isolated from swine) on the growth performance, intestinal morphology, immune-related cytokines, and gut microbiota of mice.

MATERIALS AND METHODS

Probiotic Strains and Animal Experiments

The B. velezensis JT3-1 used in this experiment was obtained from feces of healthy domestic yak (Bos grunniens) in Gansu province of China. Two strains of Lactobacillus were also isolated from healthy animal feces in Northwest China and included L. plantarum FBL-3a isolated from feces of beef cattle in Xinjiang Uygur Autonomous Region and L. salivarius ZLP-4b obtained from feces of swine in Gansu province. Our previous results found that these probiotics exhibited good tolerance and antimicrobial activity. The antibiotic sensitivity experiments, resistance gene tests, and hemolytic experiments conducted showed the safety of the JT3-1, FBL-3a, and ZLP-4b strains. Based on this, the complete genome sequence of B. velezensis JT3-1 was performed using a PacBio Sequel sequencing platform at Beijing Genewiz Bioinformatics Technology Co., Ltd.

Production Performance Analysis

In this test, the body weight of all mice were recorded at the same time every day and under identical conditions. Main organs such as the heart, lung, liver, kidney, thymus, and spleen were weighed by an electronic balance after euthanasia. The weight/body weight of an organ was expressed as organ index.
Histological Staining
The duodenum, jejunum, ileum, and colon tissues were dipped in 4% paraformaldehyde solution (Servicebio Co., Ltd., Wuhan, China) and stained using hematoxylin and eosin (H&E) solution. Images of morphological character were captured at ×40 and ×100 magnification by using a scanning electron microscope SU8100 (Hitachi Co., Ltd., Japan). The intestinal villus height, crypt depth, and the rate of villus height to crypt depth were calculated by CaseViewer 2.0 software.

Cytokine Profiling Assay
Essential cytokines in the serum of mice such as IL-4, IL-6, IL-10, IL-17A, and TNF-α were determined with Mouse ProcartaPlex Panel (Thermo Fisher Scientific, Waltham, USA) following the instructions from the manufacturer. The samples were read by Luminex TM 100/200TM instrument (Luminex Corp, Austin, TX, USA).

Fecal Extraction and 16S Sequence Analysis
Fresh feces in colon were collected under sterile conditions. The feces were promptly frozen in liquid nitrogen and then preserved at −80°C for the next step. The bacterial genomic DNA was extracted from the stored fecal pellets using QIAamp Fast DNA Stool Mini Kit (Qiagen, Germantown, MD) according to the manufacturer. The V3–V4 region of the bacterial 16S rRNA gene was PCR-amplified using the forward primer 5′-CCTACGGGNGGCWGCAG-3′ and reverse primer 5′-GACTACHVGGGTATCTAATCC-3′. The PCR product purification and amplicon library construction were performed by the Illumina NovaSeq PE250 platform at a commercial company (LC-Bio Technology Co., Ltd, Hang Zhou, China) according to the standard protocols.

Statistical Analysis
Figures were generated using GraphPad Prism 6.0 software. The statistical significance in our study was determined by t-test and χ² test by the SPSS statistical program, and P < 0.05 was considered statistically significant. The analysis was repeated for three times independently.

RESULTS
Production Performance and Organ Index Analysis
During the experiment period, no aberrant behavior was found. The overall performance in all three groups of the mice that received probiotics was improved compared to the control group. As shown in Figure 1A, there was no difference in the body weight on days 7 and 14 in the experimental and control groups, except that the weight of the mice treated with ZLP-4b was significantly increased than that of the control group on day 14 (Figure 1A). No significant difference was observed in the weight of the heart, liver, spleen, lung, thymus, and kidney between the three probiotic-administrated groups and the control group (Figure 1B). The results indicated that probiotic JT3-1, FBL-3a, and ZLP-4b administration did not affect the growth performance of mice.

Probiotic Supplementations Improve the Intestinal Mucosa Morphology
This study was aimed to investigate whether these isolated strains contribute to the intestinal mucosa function of mice. It was found that the three probiotic administration groups significantly improved the intestinal epithelial mucosal integrity in the jejunum, ileum, and colon tissues but not the duodenum compared to the control mice (Figures 2A–D). Meanwhile, the villus heights and the ratios of villi heights to crypt depths in these tissues were also increased (Figures 2E–G). These results showed that the three probiotic-administrated groups presented an improved intestinal mucosal integrity.

Effects of Probiotics on Cytokine Modulation
The serum anti-inflammatory cytokines (IL-4 and IL-10) and pro-inflammatory cytokines (IL-6, IL-17A, and TNF-α) were
FIGURE 2 | The effects of probiotic pretreatment on the epithelial mucosa integrity of the intestine of mice. (A–D) Intestinal morphology shown by hematoxylin and eosin staining of the duodenum, jejunum, ileum, and colon tissues of mice. The images of intestinal morphology at ×40 and ×100 magnification are shown, respectively. (E) Statistical analysis of villus height (µm), (F) crypt depth (µm), and (G) the ratios of villus height (µm) to crypt depth (µm) in the duodenum, jejunum, and ileum of mice, respectively. (A) JT3-1 group, (B) FBL-3a group, (C) ZLP-4b group, and (D) control group. *P < 0.05; **P < 0.01.

measured to indicate the inflammatory levels in mice after probiotics were used. The IL-10 and IL-4 levels were dramatically enhanced in the FBL-3a and ZLP-4b groups, except in the JT3-1-treated group (P < 0.01 or P < 0.001, Figures 3D,E). The content of IL-6 and TNF-α showed a reduction in the JT3-1-treated group compared to the control mice; a similar decrease was also found in FBL-3a and ZLP-4b-mice, although the changes were not distinct (P < 0.01, P < 0.05, or P > 0.05, Figures 3A,B). In addition, Figure 3C revealed that the content of IL-17A in sera was reduced after the mice received probiotics compared to the control group (P < 0.05, Figure 3C). It seemed that B. velezensis JT3-1 preferred to inhibit pro-inflammatory cytokines, while L. plantarum FBL-3a and L. salivarius ZLP-4b were more inclined to stimulate the production of anti-inflammatory cytokines.

Characterization of the Gut Microbiota in Mice After Probiotic Administration

The effects of probiotic administration on the intestinal microbial communities in mice were estimated by 16S rDNA sequence. The rarefaction curves showed that nearly all the bacteria species were sequenced in the feces of mice (Figure 4A). It was found that the mice possessed a lower bacterial species richness compared with the control group after the oral administration of probiotics (Figure 4B). Moreover, the detection of bacterial
phyla displayed that the relative abundance of Firmicutes was significantly increased, while the relative abundance of Proteobacteria and Bacteroidetes markedly declined in the fecal microbiota from the probiotic-treated mice (Figure 4C). Besides this, some phyla such as Deferrribacteres, Epsilonbacteraeota, Fusobacteria, and Actinbacteria showed a higher abundance, while lower Spirochaetes was observed in the probiotic-treated group (Figure 4D). At the genus level, the bacterial genera in the fecal microbiota showed that the relative abundance of Bilophila and Bacteroides was reduced, while that of Alistipes was increased in the probiotic supplementation group compared with that in the control group. Strangely, the Lactobacillus abundance in JT3-1 and ZLP-4b group was raised, but the FBL-3a-treated group was reduced (Figure 4E).

**DISCUSSION**

Antibiotics had a profound impact on bacterial infection in the therapeutic treatment of diseases (18). However, a number of serious problems also appeared subsequently as the synthesis of antibiotics in large quantity, such as the threat of bacterial resistance, antibiotic-associated diarrhea, and super-infection. In this study, we identified that three candidate probiotic strains from Northwest China improved the immune function and gastrointestinal health of mice. Firstly, no exceptional changes of body weight and organ index both in the control group and the treated groups demonstrated the safety of these three probiotics during the oral administration period. Secondly, the length of the villi and the villus length/crypt depth ratio were improved, and the integrity of the intestinal epithelial mucosa in the ileum and colon tissues was ameliorated, except the duodenum and jejunum segments, compared to the control mice after probiotic gavage. A similar result reported that probiotic supplementation enhanced the height of villus and the depth of crypt (19). Maybe it is just because microorganisms mainly colonize the back part of the intestine, particularly in the colon where microbes act as a major modulator of the immune system of the mucosa (20).

Cytokines play a considerable part in the immune modulatory and defense system of the host (21). However, not all probiotics could suppress pro-inflammatory cytokines and enhance anti-inflammatory factors—for instance, *Escherichia coli* strain Nissle 1917 had both pro- and anti-inflammatory effects because of the fact that *E. coli* Nissle 1917 also benefits from the inflamed environment, and it could account for why its anti-inflammatory effects are mild to moderate (22, 23). IL-4 and IL-10 play an important role in the immune modulatory pathway of the host as acknowledged inflammatory suppressor, and IL-10 production depends on IL-4 (24, 25). The cooperation between IL-10 and IL-4 suppressed Th1-related parameters (26). IL-4 is required for the induction of the class switch to IgG1 antibodies in cardiolipin-specific B cells (27). A previous study reported that the level of IL-4 was significantly evaluated after supplementation of a probiotic mixture of *L. paracasei* and *L. fermentum* (28). Some research data confirmed that IL-10 had a protective role in epithelial cells (29). In addition, IL-10 plays an effective role in inhibiting inflammation and pathogen clearance during *Borrelia recurrentis* infection (30), and *L. lactis* producing IL-10 could be a therapy of murine colitis (31). Similar effects were also found in our research that probiotic administration significantly enhanced IL-4 and IL-10 content in the FBL-3a and ZLP-4b groups. It suggested that FBL-3a and ZLP-4b have anti-inflammatory potential to some extent.

IL-6, TNF-α, and IL-17A are important pro-inflammatory mediators that play a crucial role in the processes of inflammation (32). IL-6 and TNF-α gene expression was upregulated in most intestinal inflammations such as IBD, colitis, and...
necrotizing enterocolitis (33–35). In addition, IL-6 exerts an anti-inflammatory role in a pancreatitis model by regulating the generation of cytokines, expression of adhesion agents, and activation of neutrophils (36). Two *B. subtilis* strains (BS1 and BS2) could decrease the content of TNF-α and IL-6 in a mice model (13). IL-23 promoted the pro-inflammatory cytokines IL-6 and IL-17 in autoimmune inflammatory disease models (37). IL-17A played a pivotal part in the development of dextran sodium sulfate-induced colitis by regulating the balance of Th17 and Treg cells (38). A recent research revealed that *L. helveticus* and *L. rhamnus* suppressed interleukin 17 transcription in *Citrobacter rodentium*-induced colitis in mice (39). In this study, we found that *B. velezensis* JT3-1 could diminish the content of TNF-α, IL-6, and IL-17A, while *L. plantarum* FBL-3a and *L. salivarius* ZLP-4b merely restrained the IL-17A level. Such results are in accordance with the previous studies, implying that probiotics obtained from livestock in Northwest China exerted a protective and salutary function in mice via modulating cytokine secretion.

Gut microbes play a substantial role in the maintenance of intestinal barrier function and modulation of the immune pathway (21), and the diversity of the gut microbiota is significant to the health of the hosts. We found that the species of bacteria in the gut decreased to a certain extent after probiotic administration, but some positive changes also emerged. The increase of Firmicutes/Bacteroidetes ratio contributes to nutrient intake and energy transformation (40).
Probiotics benefit a large number of pathogenic germs such as *Salmonella*, *E. coli*, *Vibrio cholerae*, and Helicobacter bacteria which can cause many infectious diseases (41). In Spirochaetes, many pathogens were identified as causes of many illnesses—for instance, *Leptospira* (leptospirosis), *Borrelia burgdorferi* (Lyme disease), *Treponema pallidum* (syphilis), *Treponema carateum* (pinta), *Borrelia recurrentis* (relapsing fever), and *Treponema pertenue* (yaws) (42–44). In this study, we found that the relative abundance of these two bacteria (Proteobacteria and Spirochaetes) was reduced by isolated probiotic pretreatment. *Bilophila* can activate Th1 cells to promote the production of IFN-γ, but it causes appendicitis as opportunistic pathogens (45). In our study, the abundance of *Bilophila* is reduced after probiotic supplementation. *Lactobacillus* is negatively correlated with TNF-α; maybe it is why the FBL-3a group with few *Lactobacillus* appeared to have higher levels of TNF-α than the other groups (46).

In our laboratory, before the animal experiment, many in vitro tests were conducted to assess the antibacterial activity and safety of JT3-1, FBL-3a, and ZLP-4b, including bile salt acid and tolerance tests, antibacterial tests, hemolytic activity tests, and antibiotic susceptibility assay (47–49). Our results demonstrated that *Bacillus* and *Lactobacillus* isolated from livestock in NorthWest China can confer benefits to mice by improving the intestinal mucosa integrity and modulating the cytokines linked to inflammation and immunity. Moreover, probiotic administration exerted positive effects on the gut microbiome of mice. This work supported the potential for *B. velezensis* JT3-1 and *L. salivarius* ZLP-4b to be functional probiotics based on their capacity of beneficial modulation of the intestinal morphology and immune-related cytokines. Further research should be performed before these strains are served in clinical practice.

DATA AVAILABILITY STATEMENT

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found below: https://www.ncbi.nlm.nih.gov/genbank/, CP032506; https://www.ncbi.nlm.nih.gov/genbank/, CP034694; https://www.ncbi.nlm.nih.gov/genbank/, CP062071.

ETHICS STATEMENT

The animal study was reviewed and approved by Lanzhou Veterinary Research Institute, Chinese Academy of Agricultural Sciences. Written informed consent was obtained from the owners for the participation of their animals in this study.

AUTHOR CONTRIBUTIONS

YiL and YoL designed this study and critically revised the manuscript. DJ, JiaW, HL, and XY performed the experiments and data analysis. JLI, JinW, GG, JLu, SX, and HY participated in the coordination and manuscript revision. All authors read and approved the final manuscript.

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