Dietary supplementation with *Bacillus subtilis* promotes growth performance of broilers by altering the dominant microbial community

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ABSTRACT The purpose of this study was to investigate the effects of *Bacillus subtilis* on growth performance, intestinal morphology, and cecal microbial composition of broilers. A total of 270 healthy one-day-old Arbor Acres male broiler chicks were randomly divided into 3 dietary treatment groups, with 6 replicates per group and 15 chickens per replicate. The dietary treatment groups were as follows: 1) basal diet, negative control group; 2) basal diet +250 g/t of zinc bacitracin, positive control group; and 3) basal diet +750 g/t of *B. subtilis*, *B. subtilis* group. Results of this experiment showed that compared with the negative control group, body weight at 42 d, average daily gain and European Production Efficiency Factor over the 42 d phase in the *B. subtilis* group and positive control group were significantly increased (*P* < 0.05); feed conversion rates in the *B. subtilis* group and positive control group were significantly decreased (*P* < 0.05); and average daily feed intake and mortality were not significantly different (*P* > 0.05). The villus height to crypt depth ratio in the ileum of the *B. subtilis* group was significantly higher (*P* < 0.05) than that of the negative control group. The results of cecal microflora at genus level were as follows. As compared with the negative control group, the abundance of *Blautia*, *Faecalibacterium*, *Flavonifractor*, and *Hydrogenoanaerobacterium* of the *B. subtilis* group and positive control group was significantly higher (*P* < 0.05), whereas the abundance of *Odoribacter* was significantly lower (*P* < 0.05). Moreover, abundance of the genera Romboutsia in the *B. subtilis* group was higher (*P* < 0.05) than that in the positive control group. The abundance of *Flavonifractor*, *Erysipelatoclostridium*, and *Hydrogenoanaerobacterium* were positively correlated with body weight and average daily gain by Spearman correlation analysis. In conclusion, dietary supplementation with *B. subtilis* improved growth performance of broilers which may be related to the increased abundance of *Blautia*, *Faecalibacterium*, *Flavonifractor*, *Hydrogenoanaerobacterium*, and Romboutsia, along with the decreased abundance of *Odoribacter*. In addition, the effect of *B. subtilis* was superior to zinc bacitracin in improving intestinal microbial composition of broilers. Therefore, *B. subtilis* may act as an effective antibiotic substitute in broilers.

Key words: *Bacillus subtilis*, 16S rDNA sequencing, growth performance, cecal microbiota, broiler

INTRODUCTION In the production of livestock and poultry, long-term abuse of antibiotics has led to several negative consequences, such as the residue of antibiotics in livestock and poultry products, the resistance of pathogens to antibiotics, the imbalance of normal microbial flora, etc. (Barton, 2000; Bogaard et al., 2000; Sorum and Sunde, 2001). With increasing food safety awareness and the introduction of relevant laws and regulations in various countries to control the use of antibiotics, the search for antibiotic alternatives has become a research focal point in the industry. One such group of antibiotic alternatives, probiotics, have been studied for nearly 20 yr (Rolfe, 2000). Probiotics have been used in livestock and poultry production as feed...
additives, and *Bacillus subtilis* is one of the most common probiotics (Guo et al., 2006).

*Bacillus subtilis* is a spore-forming aerobic bacterium. Its spores are metabolically dormant during feed processing and adaptable to external conditions, such as extremely low and high temperatures as well as low and high pH (Nicholson, 2002). In addition, dietary supplementation with *B. subtilis* can effectively improve growth performance, immunity, and intestinal morphology of poultry (Lee et al., 2010; Korosi et al., 2011; Jeong and Kim, 2014; Nguyen et al., 2015). This growth-promoting effect of *B. subtilis* may be due to its influence on gut microbial populations, including increasing the number of beneficial bacteria and reducing the number of certain pathogenic bacteria (Guo et al., 2006; Molnar et al., 2011; Wu et al., 2011; Jeong and Kim, 2014; Yang et al., 2016). Studies on the substitution of *B. subtilis* for antibiotics have also been reported (Cavazzoni et al., 1998; Lee et al., 2014). However, few studies have analyzed how *B. subtilis* as an antibiotics substitute changed the intestinal microbial community of broilers to improve the performance by microbial sequencing technology.

In this study, *B. subtilis* was used as an antibiotic substitution to determine its effect on growth performance of broilers. In addition, 16S rDNA sequencing was used to compare the effects of *B. subtilis* and the antibiotic, zinc bacitracin, on the intestinal microbial community of broilers to improve the performance by microbial sequencing technology.

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**MATERIALS AND METHODS**

**Experimental Design and Animal Management**

A total of 270 one-day old Arbor Acres male broiler chicks with similar weight (42.11 ± 0.10 g) were randomly divided into 3 groups, with 6 replicates per group and 15 chickens per replicate. The treatment groups were as follows: 1) basal diet, negative control group (NC); 2) basal diet + 250 g/t of zinc bacitracin, positive control group (PC); and 3) basal diet + 750 g/t of *B. subtilis, B. subtilis* group (BS). Zinc bacitracin and *B. subtilis* (LIFEGUFS-S 200) were provided by Lifecome Biochemistry Co., Ltd., and the number of viable *B. subtilis* in the raw product was between 2 × 10^{10} CFU/g and 3 × 10^{10} CFU/g. The testing period was from 1 to 42 d of age. Two feeding phase diets were utilized: starter diet from 1 to 21 d and grower diet from 22 to 42 d (Table 1). The diets were formulated to meet the nutrient requirements recommended by the National Research Council (NRC, 1994). The chickens were raised in an experimental farm in the Institute of Poultry Science, Chinese Academy of Agricultural Science. Birds had ad libitum to feed and water and were reared in wire cages (1.2 m × 0.9 m, length × width), with 23 h of illumination per day throughout the study. The animal use protocol was approved by the Animal Care and Use Committee of the Institute of Poultry Science, Chinese Academy of Agricultural Science (Yangzhou, Jiangsu, China).

**Growth Performance**

Daily, health status was observed, and the death and feed consumption of chickens were recorded. The birds were fasted 8 h and then weighed on day 21 and 42, to calculate the average body weight (BW), average daily gain (ADG), average daily feed intake (ADFI), feed conversion ratio (FCR), and European Production Efficiency Factor (EPEF). The EPEF was determined as per the formula (Slizewska et al., 2020): EPEF = [(viability (%) × body weight)/(FCR × age)] × 100.

**Sample Collection and Index Determination**

On day 42, 6 birds (1 bird per replicate) with similar weight per group were selected and euthanized by severing the jugular vein. About 2 cm of intestinal tissue from the duodenum, jejunum, and ileum were excised, emptied of chyme, and then fixed with 4% paraformaldehyde solution. The intestinal segments were dehydrated in an ascending gradient of ethanol. These samples were then cleaned in xylene, embedded in paraffin wax, processed into slices, and stained with hematoxylin and eosin. Villus height and crypt depth were measured using a positive fluorescence microscope (DM4000B, Leica Microsystems, Wetzlar, Germany), and villus height to crypt depth ratio (VCR) was calculated. Cecal chyme was collected, immersed in liquid nitrogen, and then stored at −80°C for DNA extraction and 16S rDNA amplicon sequencing analysis by Novogene Corporation (Beijing, China).

**DNA Extraction and Sequencing Library Construction**

Total genomic DNA was extracted from cecal contents of each chick using the EZNA Soil DNA kit (D5625-02, Omega Bio-Tek Inc., Norcross, GA). After extraction, DNA concentration and purity were analyzed by a NanoDrop ND-2000 spectrophotometer (NanoDrop Technologies, Wilmington, DE). The DNA was then stored at −20°C until further processing. DNA amplicons were amplified using primers for the V4 domain of bacterial 16S rRNA gene by polymerase chain reaction (Bergmann et al., 2011; Gao et al., 2017). The amplified products were extracted by electrophoresis with a 2% agarose gel, and the polymerase chain reaction products were mixed equally and purified with the GeneJET Gel Extraction Kit (Thermo Fisher Scientific, Waltham, MA). The library was constructed using the Ion Plus Fragment Library Kit (Thermo Fisher Scientific). After Qubit quantification (Qubit 2.0 fluorometer, Life Technology, Carlsbad) and library testing, the constructed library was sequenced using the IonS5XL sequencing platform at Novogene Bioinformatics Technology Co., Ltd. (Beijing, China).
Quality Filtering and Sequence Data Analysis

Based on the IonS5XL sequencing platform, a small fragment library was constructed using the single-end sequencing (Single-End) method. Clean data were obtained by cutting and filtering reads. Based on the clean data, the sequences were clustered into operational taxonomic units (OTU) with 97% identity, and then the OTU sequences and Silva132 database were used for species annotation analysis (Edgar, 2013). In accordance with species annotation, the differences in community structure among treatments were revealed by calculating alpha diversity and beta diversity. For alpha diversity measurements, the alpha diversity indexes were calculated based on the OTU using the Shannon, Simpson, and Chao1 (Chao, 1984; Chao and Lee, 1992) methods. For beta diversity measurements, beta diversity heatmap, principal coordinates analysis (PCoA) (Minchin, 1987), analysis of similarities (Chapman and Underwood, 1999), multiple response permutation procedure (O’Reilly and Jr, 1980), and permutation multivariate analysis of variance (Adonis) (Stat et al., 2013) were used to analyze the differences of community structure among different treatments. In addition, MetaStat (Edgar, 2004) and LEfSe analysis (Segata et al., 2011) were used to identify the biological differences between treatments. CCA-envfit function analysis (Yang et al., 2007) and Spearman correlation analysis (Segata et al., 2011) were carried out to obtain the growth performance factors which were significantly correlated with the change of community among treatments. Finally, the annotated results of the amplifier were correlated with the corresponding functional database, and functional prediction of the microbial community in the samples was carried out by Phylogenetic Investigation of Communities by Reconstruction of Unobserved States (PICRUSt) (Qin et al., 2012).

Statistical Analysis

All data were presented as the mean with pooled SEM values. Statistical analyses were carried out with SPSS 22.0 for windows (SPSS Inc., Chicago, IL). One-way ANOVA followed by LSD’s multiple comparison test was used to evaluate the differences among the treatment groups. A P-value less than 0.05 was considered statistically significant.

RESULTS

Growth Performance

Table 2 shows the growth performance and EPEF of broilers in each treatment group. Compared with the NC group, BW at 42 d (P = 0.004), ADG over the 42 d phase (P = 0.004) and EPEF (P = 0.012) in the BS group were significantly increased; and BW at 42nd day (P = 0.010), ADG over the 42-day phase (P = 0.010) and EPEF (P = 0.003) in the PC group were also significantly increased. The FCR in the BS group (P = 0.019) and PC group (P = 0.033) were significantly lower than in the NC group. There were no

Table 1. Composition and nutrient levels of the basal diet (air-dry basis).

| Items          | Contents          | Starter stage (1–21 d) | Grower stage (22–42 d) |
|----------------|-------------------|------------------------|------------------------|
| Ingredients (%)|                   |                        |                        |
| Corn           | 54.30             | 56.84                  |
| Soybean oil    | 3.40              | 3.98                   |
| Soybean meal (43%) | 38.12          | 35.32                  |
| Lysine hydrochloride (98%) | 0.15          | 0.16                   |
| DL -Met        | 0.25              | 0.24                   |
| CaCO₃          | 1.14              | 0.93                   |
| CaHPO₄.2H₂O    | 1.86              | 1.80                   |
| Salt           | 0.40              | 0.40                   |
| Choline chloride (50%) | 0.15          | 0.10                   |
| Vitamin premix¹ | 0.03            | 0.03                   |
| Mineral premix² | 0.20            | 0.20                   |
| Total          | 100.00            | 100.00                 |
| Nutrient levels (%)³ |       |                       |
| ME (kcal/kg)   | 2,950             | 3,020                  |
| CP             | 21.00             | 20.00                  |
| Ca             | 1.01              | 0.90                   |
| Available phosphorus | 0.45        | 0.43                   |
| DLys           | 1.15              | 1.10                   |
| DMet           | 0.50              | 0.48                   |
| DCys           | 0.29              | 0.28                   |
| DMet + DCys    | 0.86              | 0.82                   |

¹The vitamin premix provides the following per kg of diet: Vitamin A, 8,000 IU; Vitamin D₃, 1,000 IU; Vitamin E, 20 IU; Vitamin K₃, 0.50 mg; Vitamin B₁₂, 2.00 mg; Vitamin B₅, 8.00 mg; Vitamin B₆, 3.50 mg; Vitamin B₁₂, 0.01 mg; niacin, 35.00 mg; calcium pantothenate, 10.00 mg; folic acid, 0.55 mg; biotin, 0.18 mg.
²The mineral premix provides the following per kg of diet: Fe, 80.00 mg; Cu, 8.00 mg; Mn, 100.00 mg; Zn, 80.00 mg; I, 0.70 mg; Se, 0.30 mg.
³The nutrient levels were calculated values.
significant differences in ADFI and mortality during the 42-day phase among the 3 groups.

**Intestinal Morphology**

Intestinal morphology of the broilers at 42 d of age is shown in Table 3. Compared with the NC group, the VCR of the ileum in the BS group was significantly increased ($P = 0.047$), whereas the VCR of the duodenum and jejunum was not significantly different ($P > 0.05$) among the 3 groups. Moreover, there were no significant differences ($P > 0.05$) among any of the treatments for villus height and crypt depth of the duodenum, jejunum, and ileum.

**Variation in Cecal Microbiota Composition**

**Phylum Level** Appendix Table 1 shows the relative abundance of the top 10 microorganisms at the phylum level. The cecal microbiome of each group was dominated by Bacteroidetes, Firmicutes, Proteobacteria, Tenericutes, Melainabacteria, Actinobacteria, Cyano- bacteria, Acidobacteria, Gemmatimonadetes, and Chloroflexi. Among them, Bacteroidetes and Firmicutes were the most dominant bacterial groups, which together accounted for more than 80% of the total microbial community detected. By MetaStat analysis, Proteobacteria of the PC group was significantly higher ($P = 0.032$) than that of the BS group (Figure 1A and Appendix Table 1). However, there were no significant differences among treatments in the relative abundance of the other microorganisms in cecum of broilers at the phylum level.

**Family Level** Appendix Table 2 shows the relative abundance of the top 10 microorganisms at the family level. Compared with the NC group, the relative abundance of Ruminococcaceae ($P = 0.007$, $P = 0.018$) and Lachnospiraceae ($P = 0.019$, $P = 0.011$) in Firmicutes of the BS group and PC group were significantly increased (Figure 1B and Appendix Table 2). There was no significant difference in the relative abundance of the predominant microorganisms in cecum of broilers at the phylum level between the BS and PC groups.

**Genus Level** Appendix Table 3 shows the relative abundance of the top 35 microorganisms at the genus level. Compared with the NC group, the relative abundance of Faecalibacterium ($P = 0.006$), Flavonifractor ($P = 0.010$), Hydrogenoanaerobacterium ($P = 0.006$) and Blautia ($P = 0.009$) in Firmicutes, and Rikenella ($P = 0.047$) in Bacteroidetes of the BS group was significantly increased, and the relative abundance of Odoribacter ($P = 0.047$) in Bacteroidetes of the BS group; BW, body weight; EPEF, European Production Efficiency Factor; FCR, feed conversion rate; NC, negative control group; PC, positive control group.

### Table 2. Effects of Bacillus subtilis on growth performance and European Production Efficiency Factor of broilers at 42 d of age.

| Items       | NC     | PC     | BS     | SEM | P-value | NC-BS | NC-PC | PC-BS |
|-------------|--------|--------|--------|-----|---------|-------|-------|-------|
| BW (g)      | 29.49b | 30.58b | 30.75a | 20  | 0.004   | 0.010 | 0.651 |
| ADG (g)     | 69.2b  | 71.8a  | 72.2a  | 0.47| 0.004   | 0.010 | 0.651 |
| ADFI (g/day)| 113.3  | 114.6  | 114.9  | 0.63| 0.321   | 0.441 | 0.817 |
| FCR         | 1.637a | 1.595b | 1.590b | 0.0085| 0.019   | 0.033 | 0.781 |
| Mortality (%) | 4.45 | 1.11 | 3.33 | 1.344 | 0.748 | 0.342 | 0.523 |
| EPEF        | 410b   | 451a   | 444a   | 6.4 | 0.012   | 0.003 | 0.555 |

*a,bMeans in the same row with different superscripts are significantly different at $P < 0.05$. Values are expressed as means with pooled SEM values, $n = 6$.

Abbreviations: ADG, average daily weight gain; ADFI, average daily feed intake; BS, Bacillus subtilis group; NC, negative control group; PC, positive control group; SEM, standard error of the mean.

### Table 3. Effects of Bacillus subtilis on intestinal morphology of broilers at 42 d of age.

| Items       | NC     | PC     | BS     | SEM | P-value | NC-BS | NC-PC | PC-BS |
|-------------|--------|--------|--------|-----|---------|-------|-------|-------|
| Duodenum    | 235    | 216    | 246    | 12  | 0.730   | 0.560 | 0.388 |
| Villus height (µm) | 1,279  | 1,107  | 1,342  | 57  | 0.642   | 0.243 | 0.133 |
| VCR         | 5.5    | 5.3    | 5.8    | 0.33| 0.709   | 0.781 | 0.548 |
| Jejunum     | 237    | 220    | 215    | 10  | 0.439   | 0.532 | 0.855 |
| Villus height (µm) | 1,152  | 1,221  | 1,306  | 64  | 0.369   | 0.668 | 0.618 |
| VCR         | 4.9    | 5.8    | 6.3    | 0.36| 0.156   | 0.317 | 0.017 |
| Ileum       | 202    | 186    | 150    | 15  | 0.165   | 0.659 | 0.329 |
| Villus height (µm) | 944    | 885    | 941    | 40  | 0.981   | 0.577 | 0.593 |
| VCR         | 4.7b   | 5.5b   | 6.4a   | 0.35| 0.047   | 0.328 | 0.268 |

*a,bMeans in the same row with different superscripts are significantly different at $P < 0.05$. Values are expressed as means with pooled SEM values, $n = 6$.

Abbreviations: BS, Bacillus subtilis group; NC, negative control group; PC, positive control group; SEM, standard error of the mean.
A group was significantly decreased (Figure 1C and Appendix Table 3). In addition, when compared with the NC group, the relative abundance of Faecalibacterium (*P* = 0.024), Flavonifractor (*P* = 0.005), Hydrogenoanaerobacterium (*P* = 0.027), Blautia (*P* = 0.004), and Erysipelatoclostridium (*P* = 0.005) in Firmicutes, and Parasutterella (*P* = 0.003) and Bilophila (*P* = 0.049) in Proteobacteria of the PC group were significantly increased, and the relative abundance of Odoribacter (*P* = 0.031) in Bacteroidetes of the PC group was significantly decreased (Figure 1C and Appendix Table 3). In addition, when compared with the NC group, the relative abundance of Faecalibacterium (*P* = 0.024), Flavonifractor (*P* = 0.005), Hydrogenoanaerobacterium (*P* = 0.027), Blautia (*P* = 0.004), and Erysipelatoclostridium (*P* = 0.005) in Firmicutes, and Parasutterella (*P* = 0.003) and Bilophila (*P* = 0.049) in Proteobacteria of the PC group were significantly increased, and the relative abundance of Odoribacter (*P* = 0.031) in Bacteroidetes of the PC

**Figure 1.** Significantly different taxa between different groups by MetaStat analysis (*n* = 6). (A) Phylum. (B) Family. (C) Genus. Abbreviations: NC, negative control group; PC, positive control group; BS, *Bacillus subtilis* group.
group was significantly decreased (Figure 1C and Appendix Table 3). Compared with the PC group, the relative abundance of *Romboutsia* (*P* = 0.033) in *Firmicutes* and *Rikenella* (*P* = 0.035) in *Bacteroidetes* of the BS group was significantly increased (Figure 1C and Appendix Table 3).

**Diversity of Cecal Microbiota**

**Alpha Diversity** Alpha diversity among the NC, PC, and BS groups is presented in Table 4. Alpha diversity indexes were calculated based on the OTU using the Shannon, Simpson, and Chao1 methods. There was no significant difference in the indexes (including OTU, Shannon, Simpson, Chao1) of alpha diversity of microbiota in cecum of broilers at 42 d of age.

**Beta Diversity** Beta diversity was assessed by beta diversity heatmap and PCoA using the weighted UniFrac distance method. Beta diversity heatmap showed that microbiota diversity parameters were not affected by the treatments at 42 d of age (Figure 2A). Figure 2B shows PCoA of the variation among these 3 groups, and the cecal microbiota compositions of broilers were not separated. Analysis of similarities, multiple response permutation procedure, and Adonis are a series of nonparametric methods used to test the difference of community structure among groups. There was no significant difference in any of these 3 indexes of beta diversity of microbiota in cecum of broilers at 42 d of age (Table 5). LEfSe analysis was used to find the biomarkers with statistical difference among different treatments. Figure 2C showed the species with significant differences among the NC, PC, and BS groups with LDA scores >2.5. Seven specific biomarkers were present in the NC group, 5 in the PC group and 4 in the BS group.

**Microbiome Responding to Growth Performance**

The correlation between the dominant taxon of cecal microbiota at the genus level relative to growth performance of broilers was assessed by Spearman correlation analysis (Figure 3). The BW (Pr = 0.019), ADG (Pr = 0.019), FCR (Pr = 0.033), and EPEF (Pr = 0.038) screened by the CCA-env function analysis (Appendix Table 4) were the growth performance factors that had the most significant impact on the bacterial community. *Flavonifractor*, *Erysipelatoclostridium*, and *Hydrogenoanaerobacterium* in *Firmicutes* were positively correlated with BW and ADG. *Sphingomonas* in *Proteobacteria* was positively correlated with FCR, whereas *Bilophila* and *Parasutterella* in *Proteobacteria* were negatively correlated with FCR. *Unidentified_Clostridiales* in *Firmicutes* was negatively correlated with EPEF.

**Functional Prediction of Cecal Microbiota**

Cecal microbiota functional predictions due to dietary treatments were examined by PICRUSt. Principal component analysis revealed that the 3 dietary treatments clustered together, indicating that their functional compositions were similar (Figure 4A). Based on the Kyoto Encyclopedia of Genes and Genome, the abundant functional annotations of cecal microbiota were those corresponding to carbohydrate metabolism, amino acid metabolism, membrane transport, replication and repair, energy metabolism, translation, poorly characterized, nucleotide metabolism, cellular processes, and signaling and metabolism of cofactors and vitamins (Figure 4B). However, the predominant functions (top10) were not significantly different among the 3 groups (Figure 4C and Appendix Table 5).

**DISCUSSION**

**Growth Performance**

In the poultry industry, it is very important to find safe and effective antibiotic substitutes that also may provide economic benefits. Growth performance characteristics (including BW, ADG, ADFI, and FCR) are some of the most important factors used to evaluate the economic benefits of broiler production. The EPEF is a comprehensive measure of broiler production which reflects various measures of boiler performance, including BW, survival rate, FCR, production management, and so on. It is also a profitability index. The larger the index, the more profitable the birds are. In recent years, EPEF has gradually been recognized by practitioners and gradually become an important evaluation method of poultry production (Bhamare et al., 2016; Slizewska et al., 2020). Studies have shown that *B. subtilis* as a dietary additive can significantly promote the growth performance of broilers (Jeong and Kim, 2014; Park and Kim, 2014; Nguyen et al., 2015). Consistent with previous studies, our results showed that *B. subtilis* or zinc bacitracin can significantly increase BW.

| Table 4. Effects of *Bacillus subtilis* on alpha diversity of microbiota in cecum of broilers at 42 d of age. |
|---------------------------------|-------|-------|-------|------|-------|-------|-------|-------|
| Items                           | Groups | SEM   | NC-BS | NC-PC | PC-BS |
|---------------------------------|-------|-------|-------|------|-------|
| Observed_species                | 694   | 661   | 553   | 41   | 0.179 | 0.742 | 0.300 |
| Shannon                         | 5.53  | 5.71  | 5.69  | 0.071| 0.383 | 0.329 | 0.914 |
| Simpson                         | 0.903 | 0.924 | 0.928 | 0.0071| 0.155 | 0.233 | 0.802 |
| Chao1                           | 746   | 692   | 596   | 43   | 0.171 | 0.615 | 0.370 |

Abbreviations: NC, negative control group; PC, positive control group; BS, *Bacillus subtilis* group.
and ADG and reduce FCR during the first 42 d of production. The EPEF in the BS and PC groups was also increased as compared with the NC group, indicating that economic benefits were improved. However, some studies have reported that *B. subtilis* does not affect growth performance of broiler chickens (Lee et al., 2014), which may be related to the type and additive amount of *B. subtilis*. In our study, the growth promoting effect and economic benefit of *B. subtilis* on broilers was similar to that of zinc bacitracin according to the growth performance including the BW, ADG, FCR, and EPEF.

**Intestinal Morphology**

It is well known that intestinal morphology is an important indicator of intestine health, and villus height and crypt depth are the main indicators of intestinal digestion and absorption function as well as cell maturity rate, respectively (Paiva et al., 2014). Increased VCR can provide an intestinal environment conducive to digestion and absorption of nutrients (Montagne et al., 2003). Lee et al. (2010) showed that a diet supplemented with *B. subtilis* could promote the growth of intestinal epithelial cells, increase villus height of the small intestine, and improve

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**Figure 2.** Beta diversity of the microbiome residing in the cecal chyme of broilers at 42 d of age. (A) Beta diversity heatmap. (B) PCoA plot. (C) LDA distribution histogram (LDA scores > 2.5). Abbreviations: NC, negative control group; PC, positive control group; BS, *Bacillus subtilis* group. p, phylum; c, class; o, order; f, family; g, genus; s, species.
absorption of nutrients. However, in our study, villus height and crypt depth of the duodenum, jejunum, and ileum were not affected by the addition of *B. subtilis* or zinc bacitracin. Nevertheless, VCR of the ileum in the BS group was higher than the NC group, indicating that *B. subtilis* was beneficial to intestinal health.

**Variation in Cecal Microbiota Composition**

The composition of intestinal microflora is significant for maintaining intestinal homeostasis and health of host (Zhang et al., 2018). Cecal microflora plays an important role in chicken health and growth performance, affecting food transformation, disease resistance, and pathogen colonization ( Stanley et al., 2014; Awad et al., 2016). *Bacillus subtilis* is a kind of aerobic bacterium, which can grow in the intestinal tract and consume oxygen to maintain anaerobic environment and inhibit the growth of harmful aerobic bacteria ( Hong et al., 2005). Many studies have shown that *B. subtilis* supplementation caused a significant decrease in the numbers of *Escherichia coli* and *Salmonella*, whereas the numbers of *Lactobacillus* and *Bifidobacterium* increased in the cecum ( Wu et al., 2011; Jeong and Kim, 2014; Yang et al., 2016). Based on the results of species annotation, we analyzed the changes of cecal microbial composition at phylum, family, and genus levels to partially elucidate the growth promoting mechanisms of *B. subtilis*.

At the phylum level, *Bacteroidetes* and *Firmicutes* were the dominant bacterial groups, which together accounted for more than 80% of the total microbial community detected in our study. This is consistent with previous studies in which *Bacteroidetes* and *Firmicutes* constitute most microbial communities in chickens at the phylum level, and these bacteria are known to play a role in energy production and metabolism ( Ahir et al., 2010; Oakley et al., 2014; Pandit et al., 2018). It is worth mentioning that some reports found that the dominant phylum of the cecal community is *Firmicutes* in chickens (Awad et al., 2016; Mancabelli et al., 2016), but others have reported that the dominant phylum is *Bacteroidetes* (Mold et al., 2015; Pandit et al., 2018). The results of our study showed that *Bacteroidetes* is the dominant phylum of the cecal community in 42-day-old broilers in our trial. However, the dominant phylum may change due to the age, breed, and regional differences of selected chickens. What we assessed was final colonization of broiler chickens in the later stage of the production cycle. In addition, the abundance of *Proteobacteria* caused by supplemental zinc bacitracin was higher than that of supplemental *B. subtilis* which is similar to the previous reports (Salaeen et al., 2017; Hu et al., 2020), that is, the absorption of nutrients. However, in our study, villus height and crypt depth of the duodenum, jejunum, and ileum were not affected by the addition of *B. subtilis* or zinc bacitracin. Nevertheless, VCR of the ileum in the BS group was higher than the NC group, indicating that *B. subtilis* was beneficial to intestinal health.

**Table 5.** Anosim, MRPP and Adonis P-values based on microbial community between treatment groups.

| VS. groups | Anosim P-value | MRPP P-value | Adonis P-value |
|------------|----------------|--------------|----------------|
| NC-BS      | 0.311          | 0.377        | 0.320          |
| NC-PC      | 0.311          | 0.446        | 0.500          |
| PC-BS      | 0.968          | 0.936        | 0.882          |

Abbreviations: Adonis, permutation multivariate analysis of variance; Anosim, analysis of similarities; BS, Bacillus subtilis group; MRPP, multiple response permutation procedure; NC, negative control group; PC, positive control group.
addition of antibiotics increased the abundance of *Proteobacteria*. Salaheen et al. (2017) and Hu et al. (2020) also reported that supplementing broilers with antibiotic growth promoters (tylosin, neomycin sulfate, bacitracin, erythromycin, and oxytetracycline or virginiamycin) increased the abundance of *Proteobacteria*. It is important to note that *Proteobacteria* include some zoonotic pathogens, such as *Escherichia*, *Salmonella*, *Campylobacter*, and other notable pathogenic genera (Salaheen et al., 2017; Clavijo and Florez, 2018).

Figure 4. Impact of dietary treatment on cecal microbiota functional predictions by PICRUSt. (A) PCA plot. (B) KEGG pathway annotation. (C) The predominant functions (top10) of cecal microbiota based on KEGG. Abbreviations: KEGG, Kyoto Encyclopedia of Genes and Genome; NC, negative control group; PC, positive control group; BS, *Bacillus subtilis* group.
At the family level, the abundance of Ruminococcaceae and Lachnospiraceae in Firmicutes was influenced by addition of B. subtilis or zinc bacitracin in our study. Lachnospiraceae is a microorganism producing n-butyric acid in the intestine, which may be related to host energy regulation and intestinal mucosal integrity (Liu et al., 2018). Ruminococcaceae and Lachnospiraceae contain the genera producing short-chain fatty acids (SCFA) (Nava and Stappenbeck, 2011; Zeng et al., 2019), and SCFA could inhibit the growth and reproduction of enteropathogenic bacteria by affecting intestinal pH value, which may ultimately yield good growth performance in broilers. Therefore, the increased abundance of Ruminococcaceae and Lachnospiraceae in the BS and PC groups may be related to the improvement of ADG in broilers. This finding is supported by Ma et al. (2018) who also reported that the increased abundance of Ruminococcaceae due to B. subtilis addition was associated with increased ADG and BW.

At the genus level, the dominant bacteria species of the gut were reshaped by B. subtilis or zinc bacitracin addition in the present study. The genus Blautia is a gram-positive bacterium, which can degrade different types of carbohydrates to produce metabolites such as acetic acid and lactic acid (Liu et al., 2008), and Faecalibacterium is an important butyrate-producing bacterium in the chicken cecum (Duncan et al., 2002). Both of these bacteria can provide energy for the body and reduce inflammation, and their increased abundance is indicative of intestinal health of the host (Biddle et al., 2013; Yang et al., 2016; Abaidullah et al., 2019). Therefore, the improved growth performance of the BS and PC groups in our study may be related to the increased abundance of Blautia and Faecalibacterium, which are beneficial to intestinal health. Flavonifractor belongs to the family Ruminococcaceae in Firmicutes and contributes to butyrate production (Meng et al., 2019). It is well known that adding butyrate in the animal diet is beneficial in improving feed conversion efficiency and growth performance. Hydrogenoanaerobacterium is a proteolytic bacterium that produces sulfides through the degradation of sulfur-containing amino acids and can break the aromatic ring of plant compounds to produce SCFA (Li et al., 2017). In addition, Hydrogenoanaerobacterium has been reported to be closely related to obesity phenotypes (Jung et al., 2016). In the present study, Spearman correlation analysis showed Flavonifractor and Hydrogenoanaerobacterium in Firmicutes were positively correlated with BW and ADG. Therefore, the improved growth performance of broilers in the BS and PC groups may be due to the increased abundance of Flavonifractor and Hydrogenoanaerobacterium. Liu et al. (2019) reported that proteoglycan induced mice with Ankylosing spondylitis exhibited notably increased relative abundances of Odoribacter, and Han et al. (2020) also thought Odoribacter was positively correlated with the inflammatory state. In our study, addition of B. subtilis or zinc bacitracin both reduced the abundance of Odoribacter, but there was no difference between the PC and BS groups. Taken together, the increased abundances of Blautia, Faecalibacterium, Flavonifractor, and Hydrogenoanaerobacterium, along with the decreased abundance of Odoribacter in the gut of the BS and PC groups could have contributed to the improved growth performance of the broilers.

Erysipelatoclostridium belongs to the family Erysipelotrichaceae in Firmicutes. It has been reported that high abundance of Erysipelatoclostridium may be related to reducing feed/egg ratio of laying hens (Guo et al., 2018), but Erysipelatoclostridium is also considered to be an opportunistic pathogen (Zhao et al., 2019), which may be associated with diseases such as metabolic syndrome and gout (Smith et al., 2016; Shao et al., 2017). In the present study, the genera Parasutterella and Bilophila in Proteobacteria were very abundant in cecum of broilers in the PC group, which were significantly negatively correlated with FCR. In addition, the genera Parasutterella and Bilophila have been reported to be associated with intestinal inflammation and injury (Chen et al., 2018; Cheng et al., 2018). In our study, zinc bacitracin did not exert a positive inhibiting effect on Erysipelatoclostridium, Parasutterella, or Bilophila. Although the genera Blautia, Faecalibacterium, Flavonifractor, and Hydrogenoanaerobacterium in the PC group may have yielded improved growth performance of broilers, there is also a potential risk of certain diseases due to the enrichment of the genera Erysipelatoclostridium, Parasutterella and Bilophila in the intestine of the PC group. In addition, the genus Romboutsia is a valuable intestinal biomarker because it plays a key role in maintaining health of the host (Mangifesta et al., 2018). In our study, Romboutsia of the BS group was higher than that of the PC group, further indicating that B. subtilis was superior to zinc bacitracin in improving intestinal microbial composition in broilers.

Diversity of Cecal Microbiota

The diversity of gut microbiota is important for maintaining gastrointestinal homeostasis and is beneficial to host health (Zhang et al., 2018). In our study, dietary treatments yielded several changes to the cecal microbial composition of broilers, and LEfSe analysis further identified the species with significant differences among treatments. However, B. subtilis or zinc bacitracin addition failed to modify the overall diversity of cecal microbiota at 42 d of age. These results are consistent with previous reports. Ma et al. (2018) reported that supplemental B. subtilis also did not affect the diversity of cecal microbiota in broilers. In addition, and even more like the results of the present study, Pedroso et al. (2006) described that dietary bacitracin had no significant effect on overall microbial diversity but did alter the composition of intestinal bacterial microbiota in chickens. However, Li et al. (2019) reported that B. subtilis addition improved the diversity of jejunal microbiota at 21 d but had little effect by 42 d of age. It may be that the dynamic diversity of intestinal microbiota, which is a very complex ecosystem, shifts with the change of diet and age (Isaacson and Kim, 2012). In fact, Ballou et al. (2016) suggested microbiota were
affected more by age than treatment. In addition, dietary B. subtilis can improve overall microbial diversity of chickens infected by pathogenic bacteria such as Salmonella (Oh et al., 2017; Khan and Choussalkar, 2020). Therefore, it is possible that B. subtilis can restore microbial diversity in chickens infected by pathogenic bacteria but have little effect on overall microbial diversity of healthy chickens. However, as indicated in the previous section, B. subtilis did affect the abundance of some intestinal microorganisms.

**Functional Prediction of Cecal Microbiota**

The PICRUSt analysis was used to infer the effect of B. subtilis or zinc bacitracin on the metabolic pathways of cecal microbiota in broilers. Based on Kyoto Encyclopedia of Genes and Genome prediction, the abundant functional annotations of cecal microbiota were those corresponding to carbohydrate metabolism, amino acid metabolism, membrane transport, replication and repair, energy metabolism, translation, poorly characterized, nucleotide metabolism, cellular processes and signaling, and metabolism of cofactors and vitamins. These predicted functions of cecal microbiota in broilers were similar to those predicted by other studies (Ma et al., 2018; Hu et al., 2020). However, the addition of B. subtilis or zinc bacitracin had little effect on the predicted functions of cecal microflora in broilers. Consistent with previous studies, Ma et al. (2018) also reported that the addition of B. subtilis DSM 32315 exerted little impact on the predicted functions of cecal microbiota in broilers. These results suggest that the functional abilities of cecal microflora are stable in broilers.

**CONCLUSION**

In conclusion, dietary supplementation with B. subtilis improved growth performance including the BW, ADG, FCR, and EPEF of broilers which may be related to the increased abundance of Blautia, Faecalibacterium, Flavonifractor, Hydrogenoanaerobacterium, and Romboutsia, and the decreased abundance of Odoribacter. Moreover, the effect of B. subtilis may be superior to zinc bacitracin in improving intestinal microbial composition of broilers, which may be related to the increased abundance of Romboutsia that plays a key role in maintaining health of host.

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**DISCLOSURES**

The authors declare that they have no conflicts of interest. The authors declare that they have no financial and personal relationships with other people or organizations that can inappropriately influence the work, and there is no professional or other personal interest of any nature or kind in any product, service and/or company that could be construed as influencing the content of this article.

**SUPPLEMENTARY DATA**

Supplementary data associated with this article can be found in the online version at https://doi.org/10.1016/j.psj.2020.12.032.

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