Effects of dietary supplement with a Chinese herbal mixture on growth performance, antioxidant capacity, and gut microbiota in weaned pigs

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Weaning stress decreases the growth performance of piglets and is one of the main concerns of pig industries. Traditional Chinese herbal medicines have been used to reduce the adverse effects of weaning stress as both nutritional supplements and antibiotic substitutes. This study aimed to evaluate the effects of a Chinese herbal mixture (Kangtaile, which contained Paeonia lactiflora, licorice, dandelion, and tea polyphenols) on the growth performances, immune response, antioxidant capacity, and intestinal microbiota of weaned pigs. A total of 400 weaned pigs [Duroc × (Landrace × Yorkshire)] were randomly allocated into one of four treatments: the CON group, fed with basic diet; the HM/one.tnum group, fed with basal diet supplemented with /zero.tnum./five.tnum g herbal mixture/kg diet; the HM/two.tnum group, fed with basal diet supplemented with /one.tnum./zero.tnum g herbal mixture/kg diet; or the HM/three.tnum group, fed with basal diet supplemented with /one.tnum./five.tnum g herbal mixture/kg diet. The results revealed that dietary supplementation with the herbal mixture for /two.tnum/eight.tnum days improved average daily gain and feed conversion ratio, while decreased the diarrhea rate of weaned pigs. Moreover, dietary supplementation with the herbal mixture improved the antioxidant capacity through increasing the activity of catalase (CAT) and the total antioxidant capacity (T-AOC) level, while decreasing the concentration of malondialdehyde (MDA) in the serum. Pigs supplemented with herbal mixture presented an increased serum immunoglobulin (Ig)M level on day /one.tnum/four.tnum compared with control pigs. The herbal mixture altered the composition of intestinal microbiota by influencing the relative abundances of Firmicutes and Bacteroidetes at the phylum level. The relative abundances of the Firmicutes and Bacteroidetes were significantly related to the body weight gain of pigs. In conclusion, supplementation of herbal mixture to the
Introduction

Weaning stress often leads to diarrhea, respiratory diseases, and loss of weight gain in weaned pigs (1). Antibiotic additives were thus used to reduce weaning stress in the feed of weaned pigs (2). However, the abuse of antibiotic drugs led to antibiotic residues in animal derived foods, while antibiotic-resistant bacteria were transmitted to humans through the food chain, which led to serious health hazards for humans (3). Therefore, with the ban on antibiotics as feed additives, it is urgent to identify the safe alternatives to antibiotics to enhance the protection of both animal derived foods and human health.

In livestock production, Chinese herbal medicines have been considered as safe feed additives to substitute antibiotics in animal feed (4, 5). Because Chinese herbal medicines are natural products, they contain a variety of beneficial ingredients that maintain the overall health of animals and prevent diseases in animals (6). A previous study found that Paoniflorin is an effective active ingredient in Paeoniae Radix Alba (PRA) (dried root of Paeonia lactiflora Pall) and had anti-inflammatory and immune regulation functions (7). Dietary supplementation with peony pollen improved the growth performance, digestive capacity, catalase, and total antioxidant capacity and reduced malondialdehyde level in carps (8). Licorice has anti-inflammatory and antiviral properties, and its main effective active ingredient is glycyrrhizin (9). Previous studies have revealed that the supplementation of seaweed and licorice increased the concentration of IgA and decreased the mRNA expression level of tumor necrosis factor-α in the saliva of pigs (10). The main active ingredients of dandelion are sesquiterpene lactone and phenylpropanoid polysaccharide, which have anti-inflammatory and immune regulating effects (11). A combination of 5 g/kg garlic and 50 g/kg dandelion diet improved growth performance, immunity, and antioxidant capacity and modified the composition of intestinal microbiota in weaning pigs. This study provided new insights into the nutritional regulation effects of the herbal mixtures on weaned pigs.

KEYWORDS
weaned pigs, herbal mixture, growth performance, antioxidant capacity, gut microbiota
experimental data for supplementation of the herbal mixture as an alternative to reduce post-weaning stress in pigs.

Materials and methods

Preparation of the herbal mixture (Kangtaile)

A type of herbal mixture called Kangtaile was prepared by Wuxi Sanzhi Biotechnology Co., Ltd (Wuxi, China). The herbal mixture is a mixed combination of five commonly used herbal medicines with different proportions including: Paeoniae Radix Alba (42%), licorice (28%), dandelion (28%), and tea polyphenols (2%). The ratio of Paeoniae Radix Alba and licorice was revised on the basis of the traditional Chinese medicine prescription “Shaoyao Gancao Decocition” (25). In addition, the combination of licorice and dandelion has the functions of antioxidant, anti-inflammatory and antibacterial (26). They exist simultaneously as the main components in tea drinks (27). The herbs are dried, broken and crushed into powder, as well as screened through an 800-mesh sieve before use, and then mixed by proportion to obtain herbal mixture (dark brown powder), which was stored in a sealed container at room temperature before use. Total polysaccharides, total saponins, flavonoids, and polyphenols are the main antioxidant compounds (28). The compositions of total polysaccharide and total flavonoids in herbal mixtures mainly come from Paeoniae Radix Alba, licorice and dandelion. The compositions of total saponins in herbal mixtures mainly come from licorice. The compositions of total polyphenols in herbal mixtures mainly come from Paeoniae Radix Alba, dandelion, and tea polyphenols. The principal active antioxidant components (total polysaccharide, and total saponins) of herbal mixture were determined by ultra-performance liquid chromatography (UPLC) method based on a previous study (29). In brief, the UPLC analysis was performed using a UPLC-Xevo TQ-S mass spectrometer ( Waters Xevo TQ MS, Waters Corp., Milford, MA, USA). The following UPLC condition was used: the chromatographic column is a Kinetex C18 column (2.1 × 100 mm, 1.7 μm); Mobile phase A was acetonitrile and mobile phase B was 0.1% formic acid and injection volume 1 μL. A flow rate of 0.15 mL/min; column temperature: 45°C; total running time 12 min. Total polyphenols content was determined according to the Folin-Ciocalteu chromatometry method (30). Absorbance of the sample was measured with the microplate reader (Tecan, Austria GmbH, Grödig, Austria) at 760 nm. The total phenolic content was expressed as milligram of gallic acid equivalent per gram of herbal mixture. The determination of total flavonoids was performed by NaNO₂-Al(NO₃)₃-NaOH colorimetric method with rutin as a standard for estimating the content of total flavonoids (31). After determination, the concentration of total polysaccharide, total saponins, total polyphenols and total flavonoids in the herbal mixture were 70.5, and 24.8, 13.0, and 26.2 mg/g, respectively.

Experimental design, animals, and housing

This study was approved by the Animal Care and Use Committee of Nanjing Agricultural University (SYXX2017-0027, Nanjing, China). The experiments were conducted at Jiangsu Huaduo Animal Husbandry Technology Co., Ltd. (Xuzhou, China). A total of 400 post-weaned pigs [Duroc × (Landrace × Yorkshire)] (equal number of males and females) with body weight (BW) about 8.37 ± 0.11 kg and age of 28 days were blocked according to BW and sex and then randomly allocated into one of four treatments: the CON group, fed with the basal diet (Supplementary Table 1); the HM1 group, fed with the basal diet supplemented with 0.5 g herbal mixture/kg diet; the HM2 group, fed with the basal diet supplemented with 1.0 g herbal mixture/kg diet; or the HM3 group, fed with the basal diet supplemented with 1.5 g herbal mixture/kg diet. There were 4 replicates (containing 2 male and 2 female pig pens) per treatment and 25 pigs per pen. The detailed feed formula and the nutritional levels were presented in Supplementary Table 1. All experimental pigs were raised in the same environment for 28 days. All pens were thoroughly cleaned and disinfected to reduce viral and bacterial contamination before the pigs moved in. During the trial, air-exhaust fans and hot-blast heaters were equipped to automatically control the room temperature between 21 and 25°C. The pens (3.5 × 4.5 m) were equipped with slatted plastic floors, an adjustable stainless-steel feeder, and two nipple drinkers for ad libitum access to feed and water.

The growth performance of weaned pigs

The weaned pigs and feed consumed were weighed on d 0, 14, and 28 of the experiment period to calculate the growth performances: average daily gain (ADG), average daily feed intake (ADFI), and feed to gain ratio (F: G). The diarrhea rates were determined based on the diarrhea score of pigs at the end of the experiment. The feces of weaned pigs were assessed regularly from 08:00 to 09:00 every day according to the criteria: 1-solid, well-formed feces; 2-loose and shapeless feces; 3-runny feces; and 4-watery diarrhea (32). Pigs with a fecal score ≥ 3 were considered to have diarrhea; pigs with a fecal score of <3 were considered normal. Diarrhea symptoms and mortality, if any, were recorded daily for each pig during the trial. The diarrhea rate was calculated as the (number of pigs with diarrhea)/(number of pigs tested × total experiment days) × 100%.
Complete blood count examination

A total of 64 pigs (8 male and 8 female pigs per treatment) were randomly selected to collect 5 mL blood via jugular venipuncture into anticoagulant (heparin) tubes on day 14 and 28 of the experimental period. The blood was then divided equally into two parts: one for complete blood count (CBC) test and the other for determination of antioxidant indicators. A total of 16 CBC indicators were measured by an automatic biochemical analyzer (Mindray BC-5000, Mindray Medical International Co. LTD, Shenzhen, China). These CBC indicators include white blood cell number (WBC), Neutrophil number (NEU), Lymphocyte number (LYM), monocyte number (MON), eosinophils number (EOS), basophils number (BAS), Red blood cell number (RBC), Hemoglobin (HGB), Erythrocyte specific volume (HCT), Mean Erythrocyte Volume (MCV), Mean Erythrocyte Hemoglobin content (MCH), Mean Red blood cell hemoglobin concentration (MCHC), platelet count (PLT), mean platelet volume (MPV), platelet distribution width (PDW), and platelet hematocrit (PCT).

Antioxidant capacity and immunoglobulin M determination

The serum samples (n = 64) were obtained by centrifuging (4,000 × g for 10 min) at 4°C, and stored at −80°C until analysis. Six oxidative stress indicators, such as malondialdehyde (MDA), total superoxide dismutase (SOD), glutathione (GSH), glutathione peroxidase (GSH-Px), catalase (CAT), and total antioxidant capacity (T-AOC) were determined using ELISA kits (Nanjing Jiancheng Institute of Biological Engineering, Nanjing, China) according to the manufacturer's instructions. The sensitivities of the kits are as follows: MDA: 0.1 nmol/mL; SOD: 0.1 U/mL; GSH: 10 μmol/L; GSH-Px: 0.2 mg/mL; CAT: 0.2 U/mL; T-AOC: 0.1 mmol/gprot. The intra- and inter-assay coefficients of variation were <10 and 15%, respectively. Concentration of immunoglobulin M (IgM) was determined using an ELISA kit (Nanjing Jiancheng Institute of Biological Engineering, Nanjing, China) with an assay sensitivity of 0.1 mg/ml according to the manufacturer's instruction. A microplate reader (Tecan, Austria GmbH, Grödig, Austria) was used to determine all indicators. The intra- and inter-assay of the coefficient of variation were <10 and 15%, respectively.

Fecal sample collection, DNA extraction, and 16S rRNA gene sequencing

On day 14 and 28 of the experimental period, a total of 96 weaned pigs (6 male and 6 female pigs per treatment) were randomly selected to collect fresh feces using clean plastic bags into a 1.5 mL centrifugation tube and stored at −80°C until analysis. Genomic DNA of fecal samples was extracted by the hexadecyltrimethylammonium bromide (CTAB) method (33), and DNA purity was detected by agarose gel electrophoresis. The genomic DNA was used as the template. The V3–V4 region of the bacterial 16S rRNA gene was amplified with the specific primers (F: 5′-GTYGCAAGCMCCGGGTAA-3′; R: 5′-GGACTACHVGGGTWTCTAAT-3′). The PCR products were confirmed with 2% agarose gel electrophoresis, purified with AMPure XT beads (Beckman Coulter Genomics, Danvers, MA, USA), and then quantified by an Invitrogen Qubit 4.0 fluorometer (Invitrogen, Thermo Fisher Scientific, Waltham, MA, USA). For library construction, the NEBNext Ultra™ DNA Library Prep Kit (Illumina, United States) was used. After the library was qualified, the library was sequenced on an Illumina next-generation sequencing platform NovaSeq 6000 to generate the 250 bp paired-end reads. The original sequencing data were spliced and filtered to obtain clean clean reads. Then, through Divisive Amplicon Denoising Algorithm 2 (DADA2), the sequences with an abundance < 5 were filtered out to obtain the final Amplicon Sequence Variants (ASVs) (34). Each de-weighting sequence generated after DADA2 is called ASVs, which corresponds to the Operational Taxonomic Units (OTUs) representative sequence. The available data were then annotated and abundances were analyzed to reveal the species composition, and further Alpha and Beta diversity analyses were performed to explore differences in community structure.

Statistical analysis

Data on growth performances (body weight, average daily gain, average daily feed intake, feed to gain ratio), CBC test, antioxidant indicators, and relative abundance of microbiota were analyzed using the GLIMMIX procedure of SAS® software 9.4 (SAS Institute, Inc., Cary, NC, USA). The model included dietary treatment, gender, and their interaction as fixed effects and pen as random effect. The experimental units for the growth performance parameters (average daily gain, average daily feed intake, and feed to gain ratio) and diarrhea rate were the pen, while the experimental units for the antioxidant indicators, blood hematology parameters and gut microbiota was the individual pig. A partial correlation analysis between the gut microbiota and growth performance indicators, antioxidant indicators was carried out using R statistical software. Graphs were performed using Graphpad Prism 8.0 (GraphPad Software Inc., San Diego, CA, USA). The results were expressed as the mean ± standard error of the mean (SEM). All tests were considered statistically significant at p < 0.05 and tendencies at 0.05 < p < 0.10. P-values were rounded to three digits after the decimal point: p-values < 0.001 are shown as <0.001.
TABLE 1 Effects of herbal mixture on growth performance of weaned pigs.

| Items           | Dietary treatments2 | SEM3 | p-value |
|-----------------|---------------------|------|---------|
|                 | CON  | HM1 | HM2 | HM3 | Treatment | Sex | Treatment × sex |
| BW (kg)         |      |     |     |     |           |     |                  |
| Day 1           | 8.50 | 8.29 | 8.22 | 8.64 |           |     |                  |
| Day 14          | 12.78 | 12.82 | 13.07 | 12.99 |           |     |                  |
| Day 28          | 18.82 | 19.22 | 19.71 | 19.43 |           |     |                  |
| Days 1–14 ADG (g) | 369.03 | 391.04 | 410.04 | 384.86 |           |     |                  |
|                  | 362.42 | 386.94 | 409.52 | 387.59 |           |     |                  |
|                  | 377.25 | 378.33 | 381.46 | 369.46 |           |     |                  |
|                  | 1.56  | 1.48 | 1.41 | 1.46 |           |     |                  |
| Days 15–28 ADG (g) | 369.03 | 391.04 | 410.04 | 384.86 |           |     |                  |
|                  | 362.42 | 386.94 | 409.52 | 387.59 |           |     |                  |
|                  | 377.25 | 378.33 | 381.46 | 369.46 |           |     |                  |
|                  | 1.56  | 1.48 | 1.41 | 1.46 |           |     |                  |
| Diarrhea rate (%) | 6.38  | 4.50 | 4.42 | 3.50 |           |     |                  |

Least-square means ± S.E.M.

BW, body weight; ADG, average daily gain; ADFI, average daily feed intake; F: G, feed to gain ratio.

2Dietary treatments: CON, the control group, fed with the basal diet; HM1, the herbal mixture group 1, fed with the basal diet supplemented with 0.5 g herbal mixture/kg diet; HM2, the herbal mixture group 2, fed with the basal diet supplemented with 1.0 g herbal mixture/kg diet; HM3, the herbal mixture group 3, fed with the basal diet supplemented with 1.5 g herbal mixture/kg diet.

3SEM = Standard error of the mean.

Values in the same row not sharing a common superscript mean a significant difference (p < 0.05).

Results

Growth performance and diarrhea rate

The effects of the herbal mixture on the growth performance and the diarrhea rate of weaned pigs were presented in Table 1. There was no significant effect of dietary treatment, sex or treatment × sex interaction on initial body weight and body weight on day 14 (p > 0.05). On day 28 of the experiment period, the body weight of pigs in the HM1, HM2, and HM3 groups were greater than that of pigs in the CON group (19.17, 19.73, and 19.26 vs. 18.70 kg, p < 0.001). During the first period (day 1–14), dietary supplementation with herbal mixture tended to increase ADG compared to the CON group (325.52, 340.16, and 322.77 vs. 307.99 g/day, p = 0.064). ADFI in the HM3 group was less than that in the CON group (464.50 vs. 479.84 g/day, p = 0.002). Female pigs had a higher ADFI compared with male pigs (478.84 vs. 472.33 g/day, p = 0.018). Dietary supplementation with herbal mixture had a lower F: G compared with pigs offered the basal diets (1.48, 1.40, and 1.44 vs. 1.56, p = 0.004). During the second period (day 15–28), ADFI in the HM1 and HM2 groups were greater than that in the CON group (673.93 and 676.62 vs. 659.89 g/day, p = 0.009). Female pigs had a higher ADFI compared with male pigs (674.41 vs. 664.88 g/day, p = 0.007). In the male pigs, dietary supplementation with herbal mixture had a greater ADG compared with pigs offered the basal diets (674.57, 675.10, and 664.50 vs. 645.36 g/day, p = 0.015). Pigs offered herbal mixture had a lower F: G compared with the pigs offered the basal diets (1.49, 1.41, and 1.49 vs. 1.56, p < 0.001). Male pigs had a lower F: G compared with female pigs (1.46 vs. 1.51, p = 0.005). In the male pigs, F: G in the HM2 group was less than that of pigs in the CON group (1.42 vs. 1.49, p = 0.024). In the female pigs, dietary supplementation with herbal mixture had a lower F: G compared with pigs offered the basal diets (1.49, 1.41, and 1.49 vs. 1.52, p = 0.001). During the whole experiment period, dietary supplementation with herbal mixture had a greater ADG compared with pigs offered the basal diets (388.99, 409.78, and 386.22 vs. 365.74 g/day, p = 0.002). ADFI in the HM3 group was less than that in the CON group (566.32 vs. 576.59 g/day, p < 0.001). Dietary supplementation with herbal mixture had a lower F: G compared with pigs offered the basal diets (1.49, 1.41, and 1.47 vs. 1.58, p = 0.001).
TABLE 2 Effects of herbal mixture on antioxidant indicators and IgM concentrations in serum of weaned pigs.

| Items                  | Dietary treatment | Male                        | SEM | p-value          | Female                        | SEM | p-value          |
|------------------------|-------------------|-----------------------------|-----|------------------|-------------------------------|-----|------------------|
|                       |                   | CON | HM1 | HM2 | HM3 | CON | HM1 | HM2 | HM3 | Treatment | Sex | Treatment × sex |
| Day 14                |                   |     |     |     |     |     |     |     |     |           |     |                |
| SOD (U/mL)            |                   | 155.14 | 166.68 | 160.95 | 165.82 | 164.10 | 156.08 | 167.45 | 161.34 | 4.10   | 0.656 | 0.974 | 0.060 |
| CAT (U/mL)            |                   | 5.62  | 5.22  | 6.05  | 5.84  | 5.39  | 6.05  | 5.75  | 6.10  | 0.20   | 0.243 | 0.458 | 0.106 |
| GSH (µmol/L)          |                   | 58.47 | 59.82 | 64.67 | 64.92 | 62.25 | 63.74 | 58.45 | 64.20 | 2.02   | 0.239 | 0.899 | 0.052 |
| GSH-PX (U/mL)         |                   | 642.83 | 709.24 | 681.08 | 653.01 | 629.52 | 592.77 | 598.75 | 661.54 | 29.06  | 0.867 | 0.016 | 0.117 |
| T-AOC (U/mL)          |                   | 5.63  | 5.65  | 5.66  | 5.74  | 5.60  | 5.67  | 5.66  | 5.71  | 0.029  | 0.011 | 0.780 | 0.861 |
| MDA (nmol/mL)         |                   | 3.46  | 3.12  | 2.98  | 3.01  | 2.99  | 3.06  | 2.87  | 2.97  | 0.13   | 0.149 | 0.075 | 0.368 |
| IgM (mg/mL)           |                   | 1.86  | 1.89  | 1.93  | 1.89  | 1.86  | 1.93  | 1.90  | 1.92  | 0.02   | 0.026 | 0.645 | 0.187 |
| Day 28                |                   |     |     |     |     |     |     |     |     |           |     |                |
| SOD (U/mL)            |                   | 169.65 | 175.20 | 180.48 | 182.01 | 172.75 | 172.67 | 173.45 | 9.17  | 0.914  | 0.469 | 0.869 |
| CAT (U/mL)            |                   | 5.95  | 7.07  | 7.03  | 7.01  | 5.99  | 6.46  | 6.76  | 6.58  | 0.31   | 0.022 | 0.172 | 0.765 |
| GSH (µmol/L)          |                   | 63.02 | 66.51 | 66.54 | 66.98 | 65.36 | 66.06 | 64.63 | 63.97 | 2.19   | 0.805 | 0.639 | 0.640 |
| GSH-PX (mg/mL)        |                   | 650.53 | 641.22 | 622.65 | 664.21 | 646.32 | 660.04 | 675.39 | 603.67 | 26.42  | 0.925 | 0.932 | 0.250 |
| T-AOC (mg/gprot)      |                   | 4.35  | 4.75  | 4.54  | 4.57  | 4.38  | 4.61  | 4.79  | 4.66  | 0.11   | 0.018 | 0.770 | 0.401 |
| MDA (nmol/mL)         |                   | 3.54  | 3.23  | 3.17  | 2.99  | 3.45  | 3.29  | 3.06  | 2.93  | 0.15   | 0.015 | 0.679 | 0.952 |
| IgM (mg/mL)           |                   | 1.92  | 1.93  | 1.94  | 1.94  | 1.93  | 1.94  | 1.93  | 1.92  | 0.010  | 0.740 | 0.753 | 0.290 |

Least-square means ± S.E.M.  
SOD, superoxide dismutase; CAT, catalase; GSH, glutathione; GSH-PX, glutathione peroxidase; T-AOC, total antioxidant capacity; MDA, malondialdehyde; IgM, immunoglobulin M.

1 Data were the mean of 16 replicates per treatment.

2 Dietary treatments: CON, the control group, fed with the basal diet; HM1, the herbal mixture group 1, fed with the basal diet supplemented with 0.5 g herbal mixture/kg diet; HM2, the herbal mixture group 2, fed with the basal diet supplemented with 1.0 g herbal mixture/kg diet; HM3, the herbal mixture group 3, fed with the basal diet supplemented with 1.5 g herbal mixture/kg diet.

3 SEM = Standard error of the mean.  
abc Values in the same row not sharing a common superscript mean a significant difference (p < 0.05).

< 0.001). During the whole 28-day experimental period, dietary supplementation with herbal mixture had a lower diarrhea rate compared with pigs offered the basal diets (4.86, 4.46, and 3.79 vs. 6.30, p < 0.001).

Serum antioxidant indicators and IgM concentration

The effects of the herbal mixture on the antioxidant capacity of weaned pigs were presented in Table 2. There were evident tendencies of dietary treatment × sex interaction effects on the level of SOD (p = 0.060) and GSH (p = 0.052) in pigs on day 14. In the male pigs, the activity of SOD in the HM1 group was greater than that in the CON group (166.68 vs. 155.14 U/mL, p = 0.046). In the male pigs, the activity of GSH in the HM2 and HM3 group were greater than that in the CON group (64.67 vs. 58.47 µmol/L, p = 0.036, 64.92 vs. 58.47 µmol/L, p = 0.037, respectively). On day 14, male pigs had a higher activity of GSH-PX compared with female pigs (671.54 vs. 620.64 mg/mL, p = 0.016). On day 14, the level of T-AOC in the HM3 group was significantly greater than that in the CON group (5.73 vs. 5.62 U/mL, p = 0.001). The concentration of MDA in the female pigs had a tendency to decrease compare with the male pigs (2.97 vs. 3.14 nmol/mL, p = 0.075). Dietary supplementation with herbal mixture had a greater concentration of IgM compared with the pigs offered the basal diets (1.91, 1.91, and 1.91 vs. 1.86 mg/mL, p = 0.026). On day 28, the dietary supplementation with herbal mixture significantly increased the activity of CAT in weaned pigs (6.77, 6.89, and 6.80 vs. 5.97 U/mL, p = 0.022) compared with the CON group. The level of T-AOC in the HM groups was significantly greater than that in the CON group (4.68, 4.67, and 4.58 vs. 4.36 U/mL, p = 0.018). The concentration of MDA in the HM2 and HM3 group was significantly less than that in the CON group (3.12 and 2.97 vs. 3.50 nmol/mL, p = 0.015). There was no effect of treatment, sex or treatment × sex interaction on the activity of antioxidant indicators in the serum of pigs (p > 0.05).

Complete blood count test

The effects of the herbal mixture on the complete blood count of weaned pigs were presented in Supplementary Table 2.
FIGURE 1

Effects of herbal mixture on the intestinal microbiota diversity in weaned pigs on day 14 and day 28 based on alpha diversity parameters of chao1 index (A), observed OTUs (B), shannon index (C) and simpson index (D). The X-axis is the experimental period, and the Y-axis is the diversity indexes. CON, basic diet; HM1, basic diet with 0.5 g/kg of herbal mixture; HM2, basic diet with 1.0 g/kg of herbal mixture; HM3, basic diet with 1.5 g/kg of herbal mixture. Different lowercase letters in the figure indicate statistically significant differences ($p < 0.05$).

On day 14, female pigs had a higher concentration of MCHC compared with male pigs (317.91 vs. 309.97 g/L, $p = 0.008$). On day 28 of the trial, dietary supplementation with herbal mixture had greater number of WBC compared with pigs offered the basal diets (23.13, 24.32, and 23.98 vs. 22.60 × 10$^9$/L, $p < 0.001$). The number of NEU in the HM2 and HM3 groups were significantly greater than that in the CON group (12.36 and 11.98 vs. 10.17 × 10$^9$/L, $p = 0.001$). Similarly, the number of MON in the HM2 and HM3 groups were significantly greater than that in the CON group (1.97 and 2.04 vs. 1.70 × 10$^9$/L, $p < 0.001$). Dietary supplementation with herbal mixture had greater concentration of MCHC compared with pigs offered the basal diets (328.81, 319.69, and 323.81 vs. 313.31 g/L, $p < 0.001$). The number of PCT in the HM1 group was significantly less than that in the CON group (0.31 vs. 0.45%, $p = 0.025$).

Effect of herbal mixture on intestinal microbiota

To evaluate the effects of the herbal mixture on intestinal microbiota diversity, 16s rRNA gene sequencing was performed using the fresh feces of weaned pigs. A total of 7,205,863 clean reads were obtained from 96 fecal samples after quality control of the original raw reads with an average of 75,061 ± 1,233 reads per sample (Supplementary Table 3). These results indicated that the sequencing data met the criteria for further analysis.

Effect of herbal mixture on intestinal microbiota diversity of weaned pigs

As shown in Figure 1, chao1, observed OTUs, shannon, and simpson indexes were used to analyze the differences in Alpha diversity between groups. On day 14, chao1 and observed OTUs indexes of the HM1, HM2, and HM3 groups were greater than those of the CON group ($p < 0.05$). The Shannon index of the HM1 group was greater than that of the CON group ($p < 0.05$). The Simpson index in the HM2 group was less than that in the CON group ($p < 0.05$). On day 28, chao1, observed OTUs, and Shannon index of the HM1 and HM2 groups were less than those of the CON group ($p < 0.05$). The Simpson index of the HM1 group was less than that of the CON group ($p < 0.05$).
Effect of herbal mixture on intestinal microbiota composition and abundance in weaned pigs

The overall microbiota composition at the phylum and genus levels were presented in Figure 2. At the phylum level, the top 10 dominant bacterial species in the feces of weaned pigs were Firmicutes, Bacteroidata, Actinobacteriota, Proteobacteria, Euryarchaeota, Spirochaetaota, Acidobacteriota, Chloroflexi, Verrucomicrobiota, and Desulfobacterota. The fecal flora composition of the pigs was dominated by Firmicutes, accounting for 78.29%, and 75.44% on day 14, and 28, respectively. Bacteroidata accounted for 13.96%, and 15.79% on day 14 and 28, respectively (Figure 2A). At the genus level, the top 10 dominant bacterial species in the feces of weaned pigs were Lactobacillus, Bifidobacterium, Clostridium_sensu_stricto_1, Prevotella, Muribaculaceae, Blautia, Clostridia_UCG-014, Catenisphaera, Holdemanella, and Faecalibacterium. The fecal flora composition of the pigs was dominated by Lactobacillus, accounting for 9.79%, and 4.70% on day 14, and 28, respectively (Figure 2B).

When comparing the relative abundance of the four dominant phyla of the intestinal microbiota between groups (Figure 3), we found that there were significant differences.
Comparison of dominant intestinal microbiota at phylum level. Relative abundances of Firmicutes (A), Bacteroidetes (B), Actinobacteria (C), and Proteobacteria (D) among the four groups on d 18 and d 55. CON, basic diet; HM1, basic diet with 0.5 g/kg of herbal mixture; HM2, basic diet with 1.0 g/kg of herbal mixture; HM3, basic diet with 1.5 g/kg of herbal mixture. Different lowercase letters in the figure indicate statistically significant differences ($p < 0.05$).
Comparison of dominant intestinal microbiota at genus level. Relative abundances of *Lactobacillus* (A), *Blautia* (C), *Clostridium_UCG-014* (D), *Prevotella* (E), and *Bifidobacterium* (F) among the four groups on d 18 and d 35. CON, basic diet; HM1, basic diet with 0.5 g/kg of herbal mixture; HM2, basic diet with 1.0 g/kg of herbal mixture; HM3, basic diet with 1.5 g/kg of herbal mixture. Different lowercase letters in the figure indicate statistically significant differences ($p < 0.05$).

Male pigs had a higher relative abundance of *Firmicutes* (80.79 vs. 72.35%, $p = 0.009$), while had a lower relative abundance of *Bacteroidata* (12.36 vs. 19.22%, $p = 0.009$) compared with female pigs (Supplementary Figure 1). In the male pigs, the relative abundance of *Proteobacteria* in the HM1 group was less than that in the CON group (1.35 vs. 4.63%, $p = 0.025$) (Supplementary Figure 2B).

The relative abundance of the dominant genus of the intestinal microbiota between groups were presented in Figure 4. On day 14, the relative abundance of *Lactobacillus*
Linear discriminant analysis coupled with effect size (LEfSe). The LEfSe analysis identified the differentially abundant (LDA score > 4) bacterial taxa between the HM group and the CON group on day 14 (A) or on day 28 (C). A linear discriminant analysis (LDA) score higher than 4 indicates that the relative abundance of the corresponding group is higher than that of the other groups. The abscissa of the bar chart represents the LDA value, and the ordinate is the different species of the selected treatments. Cladogram showing the most discriminative bacterial clades identified by LEfSe between the HM group and CON group on day 14 (B) or on day 28 (D). LEfSe taxonomic clade: Different colors indicate that some taxa are enriched in different groups. The size of a circle is based on relative abundance. CON, basic diet; HM1, basic diet with 0.5 g/kg of herbal mixture; HM2, basic diet with 1.0 g/kg of herbal mixture; HM3, basic diet with 1.5 g/kg of herbal mixture.

and Blautia in the HM2 group was greater than that in the other groups (p < 0.05) (Figures 4A,B). The relative abundance of Clostridium and Prevotella in the HM groups was less than that in the CON groups (p < 0.05) (Figures 4C,D); The relative abundance of Bifidobacterium in the HM1 and HM3 groups were less than that in the CON group (p < 0.05) (Figure 4E). The relative abundance of Clostridia_UCG-014 in the HM1 group was greater than that in the CON group (p < 0.05) (Figure 4F).

In the male pigs, the relative abundance of Bifidobacterium in the HM2 group was greater than that in the CON group (0.29 vs. 0.11%, p = 0.008) (Supplementary Figure 3). On day 28, the relative abundance of Lactobacillus in the HM1 and HM2 groups was greater than that in the CON group (p < 0.05) (Figure 4A). The relative abundance of Clostridium in the HM1 group was greater than that in the CON group (p < 0.05) (Figure 4C). The relative abundance of Prevotella in the HM1 group was less than that in the CON group (p < 0.05) (Figure 4D). Female pigs had a higher relative abundance of Prevotella (6.36 vs. 4.38%, p = 0.014) compared with female pigs (Supplementary Figure 4).

To identify bacterial taxa that significantly differentiated among HM and CON groups, a linear discriminant analysis effect size (LEfSe) analysis (LDA score > 4) was performed (Figure 5). On day 14, Clostridia_UCG-014 (order), Clostridia_UCG-014 (genus), and Clostridia_UCG-014 (family) were more abundant in the HM1 group; Firmicutes (phylum), Lactobacillales (order), and Blautia (genus) were more abundant in the HM2 group; Lachnospiraceae (family), and Lachnospirales (order) were more abundant in HM3 group;
**Effect of herbal mixture on intestinal microbiota function of weaned pigs**

The Kyoto Encyclopedia of Genes and Genomes (KEGG) pathway enrichment analysis were presented in Supplementary Figure 5. The top 35 KEGG pathways between the HM and CON groups were selected from the results of species function annotation and abundance information, and then clustered by their function annotation information and the inter-group differences, to obtain a KEGG pathway enrichment analysis heatmap. Predicted KEGG pathways were mainly concentrated in ABC-2 type transport system ATP-Binding protein, ATP-binding cassette, subfamily B, bacterial, sucrose-6-phosphatase, putative ABC transport system permease protein, ABC-2 type transport system permease protein, RNA polymerase sigma-70 factor, ECF subfamily, and K02003 putative ABC transport system ATP-binding protein pathways.

**Correlation analysis between the differential microbial species and measured parameters**

The partial correlation analyses between the relative abundance of dominant bacteria and growth performance, serum antioxidant parameters were presented in Figure 6. These results indicated that ADG and F:G were positively associated with phyla *Firmicutes* (R = 0.610, R = 0.674, respectively, p < 0.05) (Figure 6A). Whereas, ADG was negatively associated with phyla *Bacteroidota* (R = −0.609, p < 0.05) (Figure 6A). Meanwhile, the abundance of *Firmicutes* was negatively associated with the diarrhea rate of pigs (R = −0.600, p < 0.05). Interestingly, the serum T-AOC were positively associated with genus *Lactobacillus*, *Catenisphaera*, and *Faecalibacterium* (R = 0.256, R = 0.247, R = 0.244, respectively, p < 0.05), while negatively associated with phyla *Proteobacteria* and genus *Clostridium* (R = −0.239, R = −0.399, respectively, p < 0.05) (Figure 6B). Furthermore, *Clostridia* and *Catenisphaera* had significant negative correlation with the concentration of serum IgM (R = −0.251, R = −0.222, respectively, p < 0.05) (Figure 6B).

**Discussion**

Herbal medicine and their extracts are widely used as health promoters and novel approaches to disease treatment (35). Meanwhile, herbal medicine feed additives have also been receiving increasing attention due to their ability to improve the nutrition and wellbeing of farm animals (36). Furthermore, herbal medicine as an alternative to antibiotic feed additives improved the growth performance, meat quality, and nutrient digestibility parameters of pigs, which contributed to reducing the cost of raising pigs and produced antibiotic-free pork products (37). Combined herbal mixture have been shown to be more effective in improving animal health than single herbal extracts (38). Herbal mixtures are also regarded as one of the most economical and effective additives due to their ease of preparation and low cost, so their use has increased in livestock production (39). However, the effect of the herbal mixture containing *Paenoniea Radix Alba*, *licorice*, *dandelion*, and tea polyphenols on weaned pig production is not yet known. The present study aims to investigate the effects of these herbal mixtures on the growth performance, antioxidant capacity, immune response, and intestinal microbiota of weaned pigs.

In the swine industry, newly weaned pigs simultaneously encounter psychological, physiological, and nutritional pressures, resulting in impaired the digestive and absorptive capacity, which has a negative impact on feed intake and daily gain of weaned pigs (40). It is reported that adding feed additives to the diet reduces the negative effects of weaning stress (41). We first investigated the effect of herbal mixtures as promising feed additives that are able to substitute antibiotics on the growth performance of weaned pigs. Our findings indicated that the dietary supplementation of herbal mixture significantly improved the growth performance of weaned pigs as demonstrated by several indicators, including the BW, ADG, ADFI, and F: G, which suggested that herbal mixture effectively helped pigs overcome weaning stress. Specifically, the body weight at the end of the experiment period and the average daily gain during the whole experiment period of the pigs supplemented with herbal mixture were significantly greater than those of the CON group. Among them, the HM2 group (basic diet with 1.0 g/kg of herbal mixture) had the highest BW and ADG at the end of the trial. Previous studies found that weaned pigs fed herbal mixtures (including *Codonopsis pilosula*, *Radix astragalus*, *R. isatidis*, *R. paeoniea*...
alba, and Atractylodes macrocephala) supplementation diets had positive effects on BW and ADG, which was in agreement with our results (42). We speculated that the beneficial effects of herbal mixtures on the growth performance of pigs due to a variety of nutrients and active ingredients in herbal medicines, which improved digestive tract function in pigs (43). In addition, we also observed that the feed intake of pigs in the HM1 and HM2 group were greater than that of CON group during the second experiment period, while the feed intake of pigs in the HM3 group was the lowest among the four groups during the whole experiment period. Due to herbal mixture ingredients including Paeoniae Radix Alba, dandelion, and tea polyphenols tasting bitter to mammals (44, 45), we speculated that the bitter taste of a high dose herbal mixture added to the diet might affect the appetite of pigs, resulting in a decrease in feed intake. Feed conversion efficiency is a very important indicator in pig production that determines the economic benefits of breeding. In the present study, the F: G of pigs supplemented with herbal mixture was significantly less than those in the CON group, which suggested that adding the herbal mixture to the diet
was helpful to improve economic benefits. A previous study indicated that supplementation of plant extract mixtures in the drinking water increased ADG and decreased F:G in weaned pigs (46). Supplementation of 1.0 g dandelion/kg improved ADG, feed efficiency, and nutrient digestibility of weaned pigs (47). These were basically consistent with the experimental results presented in our study. The present study demonstrated that adding herbal mixture to the diets of pigs improved the growth performance of pigs, including improving average daily gain and reducing feed to weight ratio. Among them, dietary supplemented with 1.0 g/kg herbal mixture had the best effect.

Early weaned pigs are vulnerable to diarrhea and even death due to weaning stress and immature intestinal development (48). In the present study, the diarrhea rate of pigs supplemented with herbal mixture was significantly lower than that of the CON group, and the frequency of diarrhea decreased with the increase of the herbal mixture supplementation. A previous study reported that plant derived polyphenol extracts reduced post-weaning diarrhea caused by E. coli in pigs (49). A traditional herbal medicine called “Tongxie Yaofang” containing paoniea lactiflora has been widely used to treat diarrhea-predominant irritable bowel syndrome (50). A herbal medicine compound containing licorice extract reduced diarrhea rates and improved the nutrient apparent digestibility of piglets (51). These observations were consistent with our results in the present study. The herbal mixture contains both Paeoniae Radix Alba, licorice and polyphenol extracts in this study, and their combination may be the reason for the significant reduction in diarrhea rate in weaned pigs.

Weaning stress disrupts free-radical metabolism and the antioxidative system, causing severe oxidative stress and mitochondrial dysfunction in pigs (52, 53). The serum antioxidant indicators like GSH, T-AOC, SOD, GSH-Px, and CAT were commonly used to assess pig antioxidant stress (54). The content of MDA reflects the degree of lipid peroxidation and impaired antioxidative capacity in animals (55). Next, we investigated the effects of the dietary supplementation of herbal mixture on antioxidant capacity of weaned pigs. In the present study, the concentrations of serum T-AOC and CAT in the herbal mixture groups were greater than that those in the CON group, whereas the concentration of serum MDA in herbal mixture group was lower. Single herbal medicine or herbal mixture extracts with antioxidant capacity have been demonstrated that the supplementation of herbal mixture to the diet decreased the oxidative stress level of weaned pigs, improved the antioxidant capacity of weaned pigs. This is probably due to the fact that the five herbs used in the herbal mixture in our study contain a variety of active ingredients with antioxidant effects.

Weaning stress causes immune system dysfunction and results in a negative impact on performance and health in pigs (62). IgM is the major serum immunoglobulin (63). The level of serum IgM was associated with the level of the autoimmune ability of piglets (64). White blood cells participate in the immune response of animals, and their abnormal increase indicates the presence of inflammation, so they are important clinical indicators for health status (65, 66). Both neutrophils and monocytes are immune-related white blood cells that aid in the host’s defense against invading pathogens (67). We next further investigated the effect of adding herbal medicine mixture to the diet on the immune index of weaned pigs. In the present study, the serum IgM concentrations of pigs fed with the herbal mixture increased, indicating that the antibody produced by B lymphocytes was increased, which was conducive to improving the humoral immunity of pigs. Similarly, previous research has shown that supplementation with herbal mixture (include dandelion and licorice) increased serum IgG and IgM concentrations in piglets (68). This is basically consistent with our research results. It had been reported previously that herbs, such as Radix Alba (69), licorice (70), dandelion (71), and tea polyphenols (72), have functions of immunomodulatory and resisting inflammation, which could be used as a new-type immunoenhancer. The synergistic effect of Radix Alba and licorice improved immunity and reduced inflammation (73). Furthermore, our results indicated that the treatment of the herbal mixture with 1.0 and 1.5 g/kg concentrations significantly increased the WBC, NEU, and MON counts compared with the CON group. As immune biomarkers, abnormal levels of white blood cell (WBC) counts indicate abnormal immune processes (74). Neutrophils are the most abundant white blood cells and an important part of the innate immune system (75). Monocytes are innate immune cells and play an important defensive role in the primary innate immune response (76). A previous study had demonstrated that green tea extract increased the WBC counts in finishing pigs (77). WBC concentration was greater in pigs fed an herbal mixture (including: Astragalus membranaceus, Codonopsis pilosula and allicin) than in the CON group (16). This indicated that the herbal mixture had the potential to attenuate inflammation and enhance immunity in weaned pigs. Mean corpuscular hemoglobin concentration (MCHC) is a parameter of risk for anemia in weaning pigs (78). In the present study, the mean corpuscular hemoglobin concentration in the HM groups was greater than that of in the
CON group, indicating that herbal mixture could reduce the incidence of anemia in weaned pigs by increasing corpuscular hemoglobin concentration. Based on the improvement of immunity indicators that accompanied supplementation with herbal mixture, we speculated that adding the herbal mixture to the feed could improve the swine immunity and also their general health status.

The gut microbiota is essential for gut homeostasis and host health (79). When people investigated the microbiota associated with health and diseases, they found that microbial diversity ensures the consistent immune regulation ability of microbiota, so it is more important to understand microbial diversity (80). Microbial diversity was assessed using the Chao1, observed_OTUs, Shannon, and Simpson index (81). In the present study, chao 1 and observed_OTUs in the HM groups were greater than those in the CON group on day 14 of the experiment. In contrast, chao 1, observed_OTUs, and Shannon index in the HM groups were less than those in the CON group on day 28 of the trial period. This indicates that the herbal mixture significantly improved the diversity of bacterial community in the early stage of the experiment. As animals aged, the number of microbial species in the feces of weaned pigs decreased after adding an herbal mixture to the diet. A previous study reported that the alpha diversity indices of intestinal microbiota in pigs changed with age, and they decreased at 35 days after weaning compared with 21 days after weaning (42). We speculated that the addition of herbal mixtures to the diet might inhibit pathogenic bacteria colonization in the intestine over time, and then reduce the diversity of intestinal flora.

In addition to its role in microbial diversity, our study also indicated that the addition of herbal mixtures to the diet had an impact on the abundance of intestinal flora in piglets. In this experiment, Firmicutes and Bacteroidetes were the dominant phyla in the gut microbiome of piglets, which was basically consistent with previous report (82). We also found that the HM2 group (basic diet with 1.0 g/kg of herbal mixture) significantly increased the relative abundance of Firmicutes, while decreasing the relative abundance of Bacteroidetes compared to the CON group. Meanwhile, the HM2 group had the highest BW and ADG at the end of the trial. This indicated obese pigs had a higher abundance of Firmicutes and a lower abundance of Bacteroidetes than lean pigs during the nursery period. Human and mouse studies yielded similar results (83, 84). In addition, the abundance of Firmicutes and Bacteroidetes was related to fat deposition in animals (85, 86). These results suggested that the intestinal microbiota (i.e., Firmicutes, Bacteroidetes) affected by the herbal mixture might affect the fat metabolism of the host, leading to fat deposition and body weight gain. Furthermore, the pathogenic bacteria decreased in the herbal mixture groups. For example, the relative abundance of Proteobacteria was lower in the HM2 group than in the control group on day 28. Studies had revealed that plant-derived polyphenols reduce the relative abundance of Proteobacteria in pigs, attenuating diarrhea and intestinal damage (87). Our study also found that the diarrhea rate of HM2 group was significantly lower than that of the control group. These results suggested that changes in the abundance of Proteobacteria in the intestine might disrupt the balance of microbial community structure and the health state of the host. Supplementing the diet with herbal mixtures was able to reduce the risk of intestinal diseases. Lactobacillus, as a beneficial bacteria, maintains the balance of intestinal microflora and enhances the human immune function (88). In the present study, the relative abundance of Lactobacillus in the HM1 and HM2 groups were greater than that in the CON group. Therefore, there was reason to believe that supplementing an appropriate dose of herbal mixture improved the relative abundance of Lactobacillus, which was conducive to improving host health. Interestingly, we found that Lactobacillus decreased but Bifidobacterium increased in the group with 1.5 g/kg herbal mixture. Previous studies showed that Lactobacillus and Bifidobacterium strains had different effects on host inflammation and intestinal microbial fermentation and composition (89). Lactic acid inhibits the proliferation of Lactobacillus during the fermentation process (90). Therefore, we speculate that the addition of high-dose herbal mixtures to the feed produce more lactic acid in the intestine, which inhibits the abundance of Lactobacillus in the intestine. Since Lactobacillus and Bifidobacterium are two major probiotics in the intestine (91), the abundance of Bifidobacterium could increase to maintain the balance of probiotics when the abundance of Lactobacillus decreases. The function prediction of the microbiota analysis suggested that the main pathways were protein transport, ATP binding, and sucrase, which indirectly reflected the microorganisms participate in the digestion and absorption of nutrients in the pig intestine. Our findings indicated that supplementing with herbal mixture to the diet had an inhibitory effect on pathogenic bacteria like Proteobacteria while increasing the relative abundance of beneficial bacteria like Lactobacillus and ultimately helping pigs cope with weaning stress.

When there are multiple variables, the correlation analysis between the two variables is often affected by other variables (92). In order to exclude the possible influence of controlling variables on other related variables, partial correlation analysis was used to assess the correlation between the relative abundance of dominant bacteria and measured parameters. Partial correlation analysis indicated that ADG was significantly positively correlated with the abundance of Firmicutes and negatively correlated with the abundance of Bacteroidetes. Previous reports indicated that the ADG of meat goats was linearly associated with the abundance of Firmicutes (93), which was consistent with our experimental observations. In this study, we did not find a significant correlation between body weight and the abundance of Firmicutes, but there was a positive correlation (R = 0.41). We speculated that the selection of
samples and the differences between individuals mask this correlation due to limited sample sizes. The abundance of Firmicutes and Bacteroides has the potential to be considered as a biomarker to evaluate the body weight gain performance of pigs. Due to pigs being closer to humans in genetics, anatomy, and physiology and often used as biomedical models (94), our findings also support the view of Firmicutes and Bacteroidetes to evaluate human obesity (95). Our study also found that the abundance of Lactobacillus was positively associated with the T-AOC activity in the serum. Lactobacillus has been shown to have an antioxidant effect and to reduce oxidative stress injury by lowering ROS level (96). This indicated that supplementing herbal mixtures to the diet was capable of enhancing the abundance of Lactobacillus and then improving the antioxidant capacity of pigs.

Conclusions

The current findings indicated that supplementation of the herbal mixture (including: Paeoniae Radix Alba, licorice, dandelion, and tea polyphenols) to the diet significantly improved the ADG, while reduced feed to gain ratio (F:G) of weaned pigs. Our results also demonstrated that the herbal mixture significantly improved the total antioxidant capacity of weaned pigs. In addition, we also found that the relative abundance of beneficial intestinal microbiota such as Lactobacillus was increased in the pigs supplemented with 1.0 g/kg herbal mixture to the diet. These results also demonstrated the potential effect of the herbal mixture as an alternative to antibiotics in promoting the growth of weaned pigs, improving the ability of antioxidant capacity, and maintaining the intestinal microflora balance and improving the overall health of the host.

Author contributions

QX and MC analyzed the data. QX wrote—review and editing and visualization. MC wrote—original draft and investigation. RJ provided the resources. MC, XZ, and ML conducted the animal experiments. JZ contributed to formal analysis. XC and CZ contributed to software and visualization. BZ contributed to the project administration, and funding acquisition, project supervision. All authors have read and approved the final version of the manuscript.

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Conflict of interest

Author RJ was employed by Wuxi Sanzhi Bio-Tech Co., Ltd.

The remaining authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fvets.2022.971647/full#supplementary-material
References

1. Campbell JM, Crenshaw JD, Polo J. The biological stress of early weaned piglets. J Anim Sci Biotechnol. (2013) 4:19. doi: 10.1186/2049-1891-4-19

2. Barton MD. Antibiotic use in animal feed and its impact on human health. Nutr Rev (2008) 12:279–99. doi: 10.1093/nutrit/08729106

3. Khan MZH. Recent biosensors for detection of antibiotics in animal derived food. Crit Rev Anal Chem. (2022) 52:780–90. doi: 10.1080/10400434.2020.1820072

4. Guo S, Liu L, Lei I, Qiu X, He C, Tang S, et al. Modulation of intestinal morphology and microbiota by dietary Macleaya cordata extract supplementation in Xuefeng Black-boned Chicken. Animal. (2021) 15:100399. doi: 10.1016/j.animal.2021.100399

5. Gong I, Yin H, Hou Y, Yin Y. Review: Chinese herbs as alternatives to antibiotics in feed for swine and poultry production: potential and challenges in application. Can J Anim Sci. (2014) 94:223–41. doi: 10.4141/cjas2013-144

6. Chen JS, Kang BJ, Yao K, Fu CX, Zhao YR. Effects of dietary Macleaya cordata extract on growth performance, immune responses, antioxidant capacity, and intestinal development in weaned piglets. J Anim Res Anim. (2019) 47:349–56. doi: 10.20840/09712119.2019.1636800

7. Ji Y, Wang T, Wei ZF, Lu GX, Jiang SD, Xia YF, et al. Paenomflorin, the main active constituent of Paonia lactiflora roots, attenuates bleomycin-induced pulmonary fibrosis in mice by suppressing the synthesis of type I collagen. J Ethnopharmacol. (2013) 149:825–32. doi: 10.1016/j.jep.2013.03.017

8. Ren HT, Huang Y, Lin L. Effects of dietary supplementation with peony pollen on growth, intestinal function, fillet quality and fatty acids profiles of common carp. Aquacult Nutr. (2012) 70:907–16. doi: 10.1111/j.1365-2095.2012.00509.x

9. Kitagawa I, Licorice root. A-natural sweetener and an important ingredient in Chinese medicine. Pure Appl Chem. (2002) 74:1149–98. doi: 10.1351/pac200274071119

10. Katayama M, Fukuda T, Okamura T, Suzuki E, Tamura K, Shimizu Y, et al. Effect of dietary addition of seaweed and licorice on the immune performance of pigs. Anim Sci J. (2011) 82:274–81. doi: 10.1111/j.1740-0929.2010.00826.x

11. Gonzalez-Castejon M, Viusol R, Rodríguez-Casado A. Diverse biological activities of dandelion. Nutr Rev. (2012) 70:534–47. doi: 10.1111/j.1753-4887.2012.00509.x

12. Samolińska W, Grela ER, Kowalczyk-Vasilev E, Kiczorowska B, Klebaniuk R, Hanczakowska E. Evaluation of garlic and dandelion supplementation on the growth performance, carcass traits, and fatty acid composition of growing-finishing pigs. Anim Feed Sci Technol. (2020) 259:114316. doi: 10.1016/j.anifeedtech.2019.114316

13. Khan N, Mulkhar H. Tea polyphenols for health promotion. Life Sci. (2007) 81:519–33. doi: 10.1016/j.lfs.2007.06.011

14. Hara H, Orita N, Hatano S, Ichikawa H, Hara Y, Matsumoto N, et al. Effect of tea polyphenols on fecal flora and fecal metabolic products of pigs. J Appl Anim Res. (2018) 47:349–56. doi: 10.1080/09712119.2018.1581686

15. Wang L, Zhou GB, Liu P, Song JH, Liang Y, Yan XJ, et al. Dissection of major constituents in Deng’s herbal tea granules by rapid resolution liquid chromatography coupled with mass spectrometry. J Pharm Biomed Anal. (2011) 56:928–36. doi: 10.1016/j.jpba.2011.08.005

16. Liu Z, Xie HL, Chen, I, Huang IH. An improved weighted partial least squares method squared with near infrared spectroscopy for rapid determination of multiple components and anti-oxidant activity of Pu-Erh tea. Molecules. (2018) 23:1058. doi: 10.3390/molecules23051058

17. Liu B, Piao X, Niu W, Zhang Q, Ma C, Wu T, et al. Kuijieyuan decotion improved intestinal barrier injury of ileal mucosal colitis by affecting TLR4-dependent PI3K/AKT/NF-kappaB oxidative and inflammatory signaling and gut microbiota. Front Pharmacol. (2020). 11:1036. doi: 10.3389/fphar.2020.01036

18. Maupetit A, Larbat R, Pernaci M, Andrieux A, Guentet C, Boutigny AL, et al. Defense compounds rather than nutrient availability shape aggressiveness trait variation along a leaf maturity gradient in a biotrophic plant-pest system. Front Plant Sci. (2018) 9:1394. doi: 10.3389/fpls.2018.01394

19. Xiang Q, Hu S, Ligaba-Osena A, Yang I, Tong F, Guo W. Seasonal variation in transcriptomic profiling of Tetrasigma hemylaeum fully developed tuberosus roots enriches candidate genes in essential metabolic pathways and phytotoxins signaling. Front Plant Sci. (2021) 12:659645. doi: 10.3389/fpls.2021.659645

20. Pedersen KS, Toft N. Intra- and inter-observer agreement when using a descriptive classification scale for clinical assessment of fecal consistency in growing pigs. Prev Vet Med. (2011)98:288–91. doi: 10.1016/j.prevetmed.2010.11.016

21. Abels E. Detection of DNA of plant pathogenic mycoplasmalike organisms by a polymerase chain reaction that amplifies a sequence of the 16S rRNA gene. Phytopathology. (1992) 82:828–32. doi: 10.1094/Phyto-828-828

22. Li M, Shao D, Zhou J, Gu J, Qin J, Chen W, et al. Signatures within esophageal microbiota with progression of esophageal squamous cell carcinoma. Chin J Cancer Res. (2020) 32:755–67. doi: 10.21147/issn.1000-9640.2020.06.09

23. Li FS, Weng JK. Demystifying traditional herbal medicine with modern approach. Nat Plants. (2017) 3:1109. doi: 10.1038/nplants.2017.109

24. Abdallah A, Zhang P, Zhong Q, Sun Z. Application of traditional Chinese herbal medicine by-products as dietary feed supplements and antibiotic replacements in animal production. Curr Drug Metabolism. (2019) 20:54–64. doi: 10.2174/1879640201920180523012920

25. Liu B, Xie HL, Chen, I, Huang IH. An improved weighted partial least squares method squared with near infrared spectroscopy for rapid determination of multiple components and anti-oxidant activity of Pu-Erh tea. Molecules. (2018) 23:1058. doi: 10.3390/molecules23051058

26. Chen CM, Chen WL, Hung CT, Lin TH, Lee MC, Chen JY, et al. Shaoyao Gancao Tang (SG-Tang), a formulated Chinese medicine, reduces aggregation and exerts neuroprotection in spinocerebellar ataxia type 17 (SCA17) cell and mouse models. Aging (2019) 11:986–1005. doi: 10.18632/aging.102186

27. Zhu N, Hou I, Ma G, Liu J. Network pharmacology identifies the mechanisms of action of Shaoyao Gancao decoction in the treatment of osteoarthritis. Med Sci Monit. (2019) 25:6501–73. doi: 10.12659/MSM.915821

28. Dufay S, Worsley A, Montellier A, Avanzí C, Ny Lg TF, et al. Herbal tea extracts inhibit Cytochromes P450 3A4 in vitro. J Pharmacol Pharmacoc. (2014) 66:1478–90. doi: 10.1111/jphp.12270

29. Liu B, Xie HL, Chen, I, Huang IH. An improved weighted partial least squares method squared with near infrared spectroscopy for rapid determination of multiple components and anti-oxidant activity of Pu-Erh tea. Molecules. (2018) 23:1058. doi: 10.3390/molecules23051058

30. Qin Y, Wang S, Wen Q, Xia Q, Wang S, Chen G, et al. Interactions between Ephedra sinica and Prunus armenica: from stereoselectivity to deamination as a metabolic detoxification mechanism of salicilin. Front Pharmacol. (2021) 12:744624. doi: 10.3389/fphar.2021.744624
40. Upadhyay SD, Kim IH. The impact of weaning stress on gut health and the mechanistic aspects of several feed additives contributing to improved gut health function in weaning piglets. A review. Animals (2021) 11:2418. doi: 10.3390/ani11082418

41. Martinez G, Dieguez SN, Fernandez Paggi MB, Riccio MB, Perez Gaudio DS, Rodriguez E, et al. Effect of fosfomycin, Cymara scolytiacea extract, desoxyrivulin and their combinations on intestinal health of weaned pigs. Anim Nutr. (2019) 3:586–95. doi: 10.1016/j.anina.2019.08.001

42. Li Y, Sun T, Hong Y, Qiao T, Wang Y, Li W, et al. Mixture of five fermented herbs (Zhizhuaisu Tk) alters the intestinal microbiota and promotes the growth performance in piglets. Front Microbiol. (2021) 12:725196. doi: 10.3389/fmicb.2021.725196

43. Wenk C. Herbs and botanicals as feed additives in monogastric animals. Asian J Anim Sci. (2003) 16:282–89. doi: 10.5713/ajas.2003.282

44. Bandopadhyay P, Ghosh AK, Ghosh C. Recent developments on polyphenol-protein interactions: effects on tea and coffee taste, antioxidant properties and the digestive system. Food Funct. (2012) 3:592–605. doi: 10.1039/c2fo00006g

45. Rolnik A, Olas B. The plants of the asteraceae family as agents in the protection of human health. Int J Mol Sci. (2021) 22:32009. doi: 10.3390/ijms22063009

46. Bontempo V, Jiang XR, Cheli F, Lo Verso L, Mantovani G, Vitari F, et al. Administration of a novel plant extract product via drinking water to post-weaning piglets affects performance on gut and health. Animal. (2014) 8:721–30. doi: 10.1071/S175173111400411X

47. Yan L, Zhang ZE, Park JC, Kim IH. Evaluation of Houttuynia cordata and Taraxacum officinale on growth performance, nutrient digestibility, blood characteristics, and fecal microbial shedding in diet for weaning pigs. Asian Australian J Anim Sci. (2012) 25:1449–44. doi: 10.5713/ajas.2012.12215

48. Chen L, Xu Y, Chen X, Fang C, Zhao L, Chen F. The maturing development of gut microbiota in commercial piglets during the weaning transition. Front Microbiol. (2017) 8:1688. doi: 10.3389/fmicb.2017.01688

49. Verhelst R, Schroven M, Buys N, Niewold T. Dietary polyphenols reduce diarrhea in enterotoxigenic Escherichia coli (ETEC) infected post-weaning pigs. Livestock Sci. (2014) 160:138–48. doi: 10.1016/j.livsci.2013.11.026

50. Li J, Cui H, Cai Y, Lin J, Song X, Zhou Z, et al. Tong-Xie-Yao-Fang regulates 5-HT level in diarrhea predominant irritable bowel syndrome through gut microbiota modulation. Front Pharmacol. (2018) 9:1110. doi: 10.3389/fphar.2018.01110

51. Chen J, Mao Y, Xing C, Hu R, Xu Z, Cao H, et al. Traditional Chinese medicine prescriptions decrease diarrhea rate by relieving colonic inflammation and ameliorating caecum microbiota in piglets. Evid Based Complement Alternat Med. (2020) 2020:3647525. doi: 10.1155/2020/3647525

52. Cao ST, Wang CC, Wu H, Zhang QH, Jiao LF, Hu CH. Weaning disrupts intestinal antioxidant status, impairs intestinal barrier and mitochondrial function, and triggers mitophagy in piglets. J Anim Sci. (2018) 96:1073–83. doi: 10.2527/jas.2017-08862

53. Novais AK, Deschene K, Martel-Kennes Y, Roy C, Laforest JP, Lessard M, et al. Weaning differentially affects mitochondrial function, oxidative stress, inflammation and apoptosis in normal and low birth weight piglets. PLoS ONE. (2021) 16:e0247188. doi: 10.1371/journal.pone.0247188

54. Pardo Z, Fernandez-Figares I, Lachmina M, Lara L, Nieto R, Seiquer I. Impact of heat stress on meat quality and antioxidant markers in Iberian pigs. Antioxidants. (2010) 1:1991. doi: 10.3390/antiox10121991

55. Bednarczuk-Misa I, Berdowska I, Zboc M, Misik A, Zielinski B, Placzkowska S, et al. Paraoxonase 1 deficiency and lipid peroxidation rise reflect a degree of brain atrophy and vascular impairment in dementia. Adv Clin Exp Med. (2020) 29:71–78. doi: 10.17212/acem.201911377

56. Liang X, Yamazaki K, Kamrutzammam M, Bi X, Pantheer A, Sano H. Effects of Chinese herbal medicine on plasma glucose, protein and energy metabolism in sheep. J Anim Sci Biotechnol. (2013) 4:51. doi: 10.1186/2449-1891-4-51

57. Huang P, Wang P, Xu J, Sun M, Liu X, Lin Q, et al. Fermented traditional Chinese medicine alters the intestinal microbiota composition of broiler chickens. Res Vet Sci. (2021) 135:85–94. doi: 10.1016/j.rvsc.2020.12.021

58. Jiang XR, Zhang HJ, Mantovani G, Alborali GL, Caputo JM, Savoini G, et al. Use of the PsycheMERGE network to investigate the association between depression polygenic scores and white blood cell count. JAMA Psychiatry. (2021) 78:1365–74. doi: 10.1001/jamapsychiatry.2021.2959

59. Lakschevitz FS, Hassanpour S, Rubin A, Fine N, Sun C, Glogauer M. Identification of neutrophil surface marker changes in health and inflammation using high-throughput screening flow cytometry. Exp Cell Res. (2016) 342:200–9. doi: 10.1016/j.yexcr.2016.03.007

60. Sampaio P, Moideen K, Ranganathan UD, Bethuramaik R. Monocyte subsets: phenotype and function in tuberculosis infection. Front Immunol. (2018) 9:17126. doi: 10.3389/fimmu.2018.01726

61. Md. Elias H. Dietary supplementation of green tea by-products on growth performance, meat quality, blood parameters and immunity in finishing pigs. J Med Plants Res. (2012) 6:2458–67. doi: 10.5897/JMPR11.1643

62. Antileo R, Figueroa J, Valenzuela C. Characterization of a novel encapsulated oral iron supplement to prevent iron deficiency anemia in neonatal piglets. J Anim Sci. (2016) 94:157–60. doi: 10.2527/jas.2015-9698

63. Vital M, Gao J, Rizzo M, Harrison T, Tiedje JM. Diet is a major factor governing the fecal byproduct-producing community structure across Mammalia, Aves and Reptilia. ISME J. (2015) 9:832–43. doi: 10.1038/ismej.2014.179

64. Geva-Zatorsky N, Sefik E, Kua L, Pasman L, Tan TG, Ortiz-Lopez A, et al. Mining the human gut microbiota for immunomodulatory organisms. Cell. (2017) 168:928–45.e11. doi: 10.1016/j.cell.2017.01.022

65. Miller GE, Engen PA, Gillevet PM, Shakil MK, Sarkroodi M, Forsyth CR, et al. Lower neighborhood socioeconomic status associated with reduced diversity in the gut microbiome of low birth weight infants. JAMA Psychiatry. (2021) 78:1365–74. doi: 10.1001/jamapsychiatry.2021.2959
of the colonic microbiota in healthy adults. PLoS ONE. (2016) 11:e0148952.
doi: 10.1371/journal.pone.0148952
82. Isaacson R, Kim HB. The intestinal microbiome of the pig. Anim Health Res
Rev. (2012) 13:180–9. doi: 10.1017/S1466252312000084
83. Turbabaugh PJ, Ley RE, Mahowald MA, Magrini V, Mardis ER, Gordon JL. An
obesity-associated gut microbiome with increased capacity for energy harvest.
Nature. (2006) 444:1027–31. doi: 10.1038/nature05414
84. Magne F, Gotteland M, Gauthier L, Zazueta A, Pesoa S, Navarrete P; et al. The
firmicutes/bacteroidetes ratio: a relevant marker of gut dysbiosis in obese patients?
Nutrients. (2020) 12:1474. doi: 10.3390/nu12051474
85. Turbabaugh PJ, Backhed F, Fulton L, Gordon JL. Diet-induced obesity is linked
to marked but reversible alterations in the mouse distal gut microbiome. Cell Host
Microbe. (2008) 3:213–23. doi: 10.1016/j.chom.2008.02.015
86. Wang H, Ni X, Qing X, Zeng D, Luo M, Liu L, et al. Live probiotic Lactobacillus johnsonii
BS15 promotes growth performance and lowers fat deposition by improving lipid metabolism, intestinal
development, and gut microflora in broilers. Front Microbiol. (2017) 8:1073.
doi: 10.3389/fmicb.2017.01073
87. Xu B, Qin W, Xu Y, Yang W, Chen Y, Huang J, et al. Dietary quercetin supplementation attenuates dia phrags and intestinal damage by regulating gut microbiota in weaning piglets. Oxid Med Cell Longev. (2021) 2021:6221012.
doi: 10.1155/2021/6221012
88. Hevia A, Delgado S, Sanchez B, Margolles A. Molecular players involved in the interaction between beneficial bacteria and the immune system. Front Microbiol. (2015) 6:1285. doi: 10.3389/fmicb.2015.01285
89. Wang J, Tang H, Zhang C, Zhao Y, Derrien M, Rocher E, et al. Modulation of gut microbiota during probiotic-mediated attenuation of metabolic syndrome in high fat diet-fed mice. ISME J. (2015) 9:1–15. doi: 10.1038/ismej.2014.99
90. Chen S, Niu H, Wu Y, Sun J, Han X, Zhang L. Influence of lactic acid on cell cycle progressions in Lactobacillus bulgaricus during batch culture. Appl Biochem Biotechnol. (2021) 193:912-24. doi: 10.1007/s12010-020-03459-8
91. Tankou SK, Regev K, Healy BC, Tjon E, Lathi L, Cox LM, et al. A probiotic modulates the microbiome and immunity in multiple sclerosis. Ann Neurol. (2018) 83:1147–61. doi: 10.1002/ana.25244
92. Luo ZC, Delvin E, Fraser WD, Audibert F, Deal CL, Julien P, et al. Maternal glucose tolerance in pregnancy affects fetal insulin sensitivity. Diabetes Care. (2010) 33:2055–61. doi: 10.2337/dc10-0819
93. Min BR, Gurung N, Shange R, Solaiman S. Potential role of rumen microbiota in altering average daily gain and feed efficiency in meat goats fed simple and mixed pastures using bacterial tag-encoded FLX amplicon pyrosequencing. J Anim Sci. (2019) 97:3523–34. doi: 10.1093/jas/skz193
94. Yan S, Tu Z, Liu Z, Fan N, Yang H, Yang S, et al. A huntingtin knockin pig model recapitu lates features of selective neurodegeneration in Huntington’s disease. Cell. (2018) 173:899–1002.e13. doi: 10.1016/j.cell.2018.03.005
95. Seokkianan S, Salatino A, Castano GO, Landa MS, Fijalkowcy C, Garaycochea M, et al. Intrahepatic bacterial metataxonomic signature in non-alcoholic fatty liver disease. Gut. (2020) 69:1483–91. doi: 10.1136/gutjnl-2019-318811
96. Kong Y, Olejar KJ, On SLW, Chehlicani Y. The potential of Lactobacillus spp. for modulating oxidative stress in the gastrointestinal tract. Antioxidants. (2020) 9:610. doi: 10.3390/antiox9070610