Effect of Dietary *Fructus mume* and *Scutellaria baicalensis* Georgi on the Fecal Microbiota and Its Correlation with Apparent Nutrient Digestibility in Weaned Piglets

Feng Zhang 1,2,*, Erhui Jin 1,2,3, Xiaodan Liu 1, Xu Ji 4 and Hong Hu 1

1 College of Animal Science, Anhui Science and Technology University, Chuzhou 233100, China
2 Anhui Province Key Laboratory of Animal Nutrition Regulation and Health, Chuzhou 233100, China
3 Anhui AnFengT Animal Medicine Industry Co., Ltd., Hefei 230031, China
4 Anhui Province Key Laboratory of Livestock and Poultry Product Safety Engineering, Institute of Animal Science and Veterinary Medicine, Anhui Academy of Agricultural Sciences, Hefei 230031, China

* Correspondence: zhangfeng@ahstu.edu.cn

**Simple Summary:**** Traditional Chinese medicine (TCM) is based on ancient Chinese medical principles. In China, these medicines have played a marked role in treating various diseases and maintaining human health for thousands of years. TCM is also increasingly considered a potential alternative to the use of antibiotics in pig production and has attracted a great deal of research interest because it is simple, convenient, cheap, and effective. However, there are few studies on the effects of dietary TCM supplementation on the gut microbiota and the apparent nutrient digestibility of weaned piglets. In our study, dietary *Fructus mume* and *Scutellaria baicalensis* Georgi improved growth performance and increased the apparent ether extract (EE) digestibility by modulating gut microbial composition and structure, favoring the health of weaned piglets.

**Abstract:** Traditional Chinese medicine (TCM) has long been demonstrated to exert a therapeutic effect on various diseases and has been used as a substitute for antibiotics in pig production. However, few studies have investigated the relationship between the intestinal microbiota and apparent nutrient digestibility when weaned piglet diets are supplemented with TCM. One hundred and sixty-two 25-day-old weaning piglets were housed in an environmentally controlled nursery facility and fed a basal diet (control group, $n = 54$) or a TCM complex (*Fructus mume* 1%, *Scutellaria baicalensis* Georgi 3%) (TCM group, $n = 54$), or a fermented diet with a complex of these two TCMs (F-TCM group, $n = 54$). Compared with the control group, in the TCM and F-TCM groups, the average daily gain (ADG) increased ($p < 0.05$), the F:G ratio and diarrhea rate decreased ($p < 0.05$), and the apparent digestibility of dry matter (DM) and ether extract (EE) of weaned piglets increased ($p < 0.05$). *Bacteroidetes* and *Firmicutes* were the predominant phyla, representing approximately 95% of all sequences. The abundance of four genera and 10 OTUs (belonging to *Ruminococcaceae* UCG-014, *Lachnoclostridium*, *Prevotellaceae* NK3B31 group, *Prevotella*_1) were negatively correlated with apparent EE digestibility ($p < 0.05$). The results suggest that weaned piglets fed with antibiotic-free diets supplemented with *Fructus mume* and *Scutellaria baicalensis* Georgi gained more weight and were healthier. When added to the diet, the complex of these two TCMs may have a direct impact on apparent EE digestibility by modifying the gut microbial composition, which favors the health of weaned piglets.

**Keywords:** traditional Chinese medicine; weaned piglets; apparent nutrient digestibility; gut microbiota

1. Introduction

Traditional Chinese medicine (TCM) is used under the guidance of ancient Chinese medicinal philosophies [1]. TCM has played a marked role in disease prevention and health
improvement and has been investigated for thousands of years in China [2–4]. With a desperate worldwide need to reduce antibiotic usage in human and veterinary medicine, research into the therapeutic effects of TCM has attracted much attention in recent years. TCM has been demonstrated to exert a therapeutic effect on various diseases, such as diabetes [2], hypertension [5], gastric cancer [6], ulcerative colitis [7], colorectal cancer [8], etc. Knowledge about the underlying pharmacological mechanisms of TCM is still scarce. It has been suggested that the therapeutic effect of TCM is closely related to the gut microbiota [1,9], which is the bridge between the body and the external environment, as there is a reciprocal link between the two. On the one hand, the improvement produced by the pharmacological activity of TCM depends on the gut microbiota [6,7]. The gut microbiota can promote the transformation and metabolism of TCM components by metabolizing TCM into specific molecules, such as alkaloids, flavonoids, and polysaccharides, which are easily absorbed in the intestine [2,6]. For example, paeoniflorin can be catalyzed into paeoniflorgenin and paeoniflorin by Lactobacillus brevis and Bacteroides fragilis, and puerarin is converted into daidzein by Bifidobacterium and E. faecalis, and aconitine can be decomposed into lipoaconitine by Clostridium butyricum and B. fragilis [6]. On the other hand, when present in the digestive tract, TCM can promote the growth of probiotic bacteria and inhibit pathogens, as well as prevent bacterial transmission, thus regulating the microenvironment and maintaining the balance of the microflora [6,7]. For example, flavonoids, polysaccharides, and saponins in TCM serve as prebiotics that regenerate the gut microbiota. Escherichia coli can be directly inhibited by cinnamon essential oil [8]. The omics technologies, such as microbiomics and metabolomics, have been considered pivotal tools to help us understand the underlying mechanisms between TCM and gut microbiota.

Apparent nutrient digestibility indicates the digestibility of feed ingredients by animals and is also an important indicator used to evaluate the nutritional value of feed ingredients in diets. Previous studies had reported that dietary supplementation with TCM improved the growth performance of heat-stressed beef cattle by increasing the apparent digestibility of organic matter (OM), crude protein (CP), and acid detergent fiber (ADF) when TCM plus γ-aminobutyric acid (GABA) was used in the diet, and the apparent EE and neutral detergent fiber (NDF) digestibility also increased [10,11]. Furthermore, in lambs and hogs, dietary TCM increased the apparent digestibility of DM, OM, CP, and NDF [12]. The research results cited above may suggest that TCM supplementation of the diet would be favorable for improving the apparent nutrient digestibility of feed. Similar studies on pig models have rarely been reported.

During weaning, piglets have to face a number of challenges that are critical and stressful [13]. On the one hand, to accelerate the pace of banning the use of antibiotic growth promotors (AGPs) in China, the addition of TCM to the feed has been considered a substitute for antibiotics as a feed additive [14]. On the other hand, physiological characteristics after weaning indicate that the weaned piglets may have unique intestinal microflora [15]. Thus, at this particular stage, studies on the effect of dietary TCM on the gut microbiota and the apparent nutrient digestibility of weaned piglets have an important scientific value. Studies unraveling the relationship between the two related research strands have not been reported so far.

The present study was conducted in two stages. First, TCMs with potent antibacterial properties were selected for further application in the production of fermented feed. Second, the growth performance, apparent nutrient digestibility, and fecal microbiota of weaned piglets were investigated. This study focused on the correlation between gut microbial communities and apparent nutrient digestibility in piglets.

2. Materials and Methods

2.1. Antibacterial Susceptibility Testing

The antibacterial properties of eight TCMs, namely Fructus mume, Scutellaria baicalensis Georgii, Rhizoma imperatae, Paeoniae radix alba, Plantaginis semen, Eclipta prostrata, Fructus arctii, and Portulaca oleracea L., were determined for both Escherichia coli and Salmonella
isolates. All the minimum inhibitory concentration (MIC) tests were conducted twice in order to ensure that the results were representative. The MICs of selected TCMs were determined by the agar dilution method [16,17]. The methods and standards followed the relevant regulations of the Clinical and Laboratory Standards Institute (CLSI) [18]. The double dilution method was used to dilute the eight TCM infusions to the required concentration gradients, then sterile Mueller–Hinton (MH) agar was added and mixed to prepare the agar plates, and the bacterial suspensions (Escherichia coli and Salmonella) with a turbidity of 0.5 MCF were diluted and inoculated on the MH agar plates [17] in an inverted culture for 16–18 h at 37°C. The MICs of the Chinese medicines were recorded when the plates showed no bacterial growth.

2.2. Preparation of Fermented MIXED Feed

The Bacillus subtilis, Lactobacillus, and Saccharomyces strains used in the present experiment were isolated and generously given by the College of Food Science and Technology, Guangdong Ocean University. A 300 g basal substrate including corn and soybean meal (mass ratio 3:1), as well as TCMs with potent antibacterial properties (Fructus mume or Scutellaria baicalensis Georgi in proportions of 1%, 3%, and 10%), was mixed and supplemented with sterile water to achieve a 60% moisture content. The mixed substrate was divided into two treatment parts: one part was inoculated with B. subtilis, Lactobacillus, and Saccharomyces (a proportion of 0.2% each, 10⁸ cfu/g) and then transferred to a plastic bag, while the other part of the mixed substrate was directly transferred to a plastic bag; all plastic bags were sealed and incubated at 37°C for 144 h. The composition of the fermented mixed feeds is shown in Table 1.

Table 1. The composition of fermented mixed feeds.

| No. | Mixed Substrate, g | Fermentation Strain, g |
|-----|--------------------|------------------------|
|     | Corn | Soybean | Fructus mume | Scutellaria baicalensis Georgi | Bacillus subtilis | Lactobacillus | Saccharomyces |
| 1   | 222.75 | 74.25 | 3.00 | - | 0.60 | 0.60 | 0.60 |
| 2   | 218.25 | 72.75 | 9.00 | - | 0.60 | 0.60 | 0.60 |
| 3   | 202.50 | 67.50 | 30.00 | - | 0.60 | 0.60 | 0.60 |
| 4   | 222.75 | 74.25 | - | 3.00 | 0.60 | 0.60 | 0.60 |
| 5   | 218.25 | 72.75 | - | 9.00 | 0.60 | 0.60 | 0.60 |
| 6   | 202.50 | 67.50 | - | 30.00 | 0.60 | 0.60 | 0.60 |
| 7   | 225.00 | 75.00 | - | - | 0.60 | 0.60 | 0.60 |

Mixed substrates fermented without the fermentation strains

| No. | Mixed Substrate, g | Fermentation Strain, g |
|-----|--------------------|------------------------|
| 8   | 222.75 | 74.25 | 3.00 | - | - | - |
| 9   | 218.25 | 72.75 | 9.00 | - | - | - |
| 10  | 202.50 | 67.50 | 30.00 | - | - | - |
| 11  | 222.75 | 74.25 | - | 3.00 | - | - |
| 12  | 218.25 | 72.75 | - | 9.00 | - | - |
| 13  | 202.50 | 67.50 | - | 30.00 | - | - |
| 14  | 225.00 | 75.00 | - | - | - | - |

2.3. Fermented Mixed Feed Parameters

Each of the 14 mixed fermentation treatments was prepared in 6 repeated plastic bags, corresponding to a 6-day (144 h) fermentation. The pH value was recorded every 24 h, and a digital pH meter was used to measure the pH of samples after calibration with standard buffers (pH 4.0 and 7.0). The results of pH values at different fermentation times is shown in Table S1. Miscellaneous bacteria identification and antibacterial susceptibility tests were performed every 24 h.
2.4. Animals, Diets, and Experimental Design

This study was approved by and implemented under the supervision of the guidelines for the care and use of experimental animals of the Ministry of Science and Technology of the People’s Republic of China (Approval Number: 2006-398). The experimental protocols were approved by the experimental Animal Ethical Committee of Anhui Science and Technology University.

In total, 162 crossbred (Duroc × Yorkshire) weaning piglets (weaned at 25 days of age) with an initial average body weight (BW) of 7.55 ± 0.47 kg were selected for the 30-day experiment. All the piglets were randomly assigned to 3 dietary groups with 3 replications per treatment and 18 pigs per pen. The BW and sex were balanced among each treatment as follows: (1) piglets in the control group fed the basal diet (control group; n = 54); (2) piglets in the treatment group fed a diet with added selected TCMs with potent antibacterial properties (Fructus mume 1%, and Scutellaria baicalensis Georgi 3%) added to the basal diet (TCM group; n = 54); (3) piglets in the fermentation treatment group fed a diet supplemented with a selected TCM complex (Fructus mume 1%, Scutellaria baicalensis Georgi 3%), fermentation strains (B. subtilis, Lactobacillus, and Saccharomyces; 0.2% each, 10^8 cfu/g), and sterile water, fermented at 37 °C for 144 h (F-TCM group; n = 54) (Table 2). All corn- and soybean-based diets had no antibiotics and conformed to the nutrient requirements of the US National Research Council [19]. Environmentally controlled nursery facilities with slatted plastic flooring and mechanical ventilation were used to house the pigs. The pre-feeding period of the weaned piglets was as follows: from 25 to 30 days of age, all piglets were fed with the basal diet, and the formal feeding trial was from 30 to 60 days of age. All pigs were fed twice a day individually (at 06:00 and 18:00) and allowed free access to feed and ad libitum water during the entire experimental period. There were no antibiotics in the feed or administered therapeutically to the pigs.

Table 2. Composition and nutrient levels of the experimental diets.

| Ingredient (%) | Control | TCM | F-TCM |
|----------------|---------|-----|-------|
| Ingredient (DM) |         |     |       |
| Corn           | 67.00   | 63.00 | 62.40 |
| Soybean meal   | 25.00   | 25.00 | 25.00 |
| Self-made premix | 8.00   | 8.00  | 8.00  |
| Fructus mume   | -       | 1.00 | 1.00  |
| Scutellaria baicalensis Georgi | - | 3.00 | 3.00 |
| Bacillus subtilis | -    | -    | 0.20  |
| Lactobacillus  | -       | -    | 0.20  |
| Saccharomyces  | -       | -    | 0.20  |
| Nutrition level (%) |         |     |       |
| Crude protein (CP) | 22.97 ± 0.82 | 26.75 ± 0.73 | 27.85 ± 0.82 |
| Ether extract (EE) | 16.27 ± 0.58 | 15.48 ± 0.44 | 16.00 ± 0.24 |
| Crude fiber (CF) | 5.26 ± 1.57 | 6.29 ± 1.79 | 7.15 ± 0.46 |
| Ash            | 5.40 ± 0.00 | 4.54 ± 0.02 | 5.36 ± 0.27 |
| Nitrogen-free extract (NFE) | 37.36 ± 2.99 | 44.21 ± 3.22 | 38.99 ± 4.59 |

Table 2. Composition and nutrient levels of the experimental diets.

*Ingredients: fish meal, choline chloride, vitamin, mineral elements, L-lysine hydrochloride, calcium hydrophosphate, stone powder, sodium chloride, enzyme preparation, flavoring agent, and sweetening agent. No antibiotics were added. Measured values (Mean ± SD).*

2.5. Determination of the Growth Performance and Diarrhea Rate

The individual BW of pigs with empty stomachs was measured at 05:00 on Days 1, 15, and 30 of the experimental period; the feed offered and residual feed were weighed and recorded daily and used to determine the ADG, average daily feed intake (ADFI), and feed-to-gain (F:G) [11]. The incidence and severity of piglet diarrhea were assessed by fecal consistency. If the piglets had moderately fluid feces and frothy diarrhea, in which the feces were definitely unformed and very watery, they were considered to be diarrheic [20]. The
diarrhea rate of piglets was recorded daily and calculated as follows: diarrhea rate (%) = the number of pigs with diarrhea × diarrhea days/(total number of pigs × experimental days) × 100%, where the number of pigs with diarrhea was defined as the number of piglets with diarrhea observed each day [21].

2.6. Assessment of Apparent Nutrient Digestibility

According to the method described by Fouhse et al. [22] and Niu et al. [15], samples of feed were collected daily during the experimental period, and the total daily feces from each pig were collected on the last 3 days. The collected feces for each piglet were composited and mixed thoroughly; approximately 100 g of feces was subsampled after thorough mixing, and all feed and feces samples were dried at 65 °C to constant weight for subsequent analysis.

Acid-insoluble ash (AIA) was used as an indigestible marker to assess the digestibility of the dietary components according to the procedure of the Association of Official Analytical Chemists (AOAC 942.05) [23]. The DM was analyzed according to the procedures described by Xie et al. [24]. The EE content was measured using the Soxhlet extraction method (AOAC 920.85), which was performed with a Soxhlet apparatus [15]. The CP content was measured via the Kjeldahl method (AOAC 984.13) using a Kjeltec 8400 analyzer unit (Foss, Beijing, China). CF analysis was carried out using the ANKOM A200 filter bag technique (AOAC 962.09) [15]. The ingredient composition and nutrient specifications of the basal and experimental diets were calculated as previously reported, and the apparent nutrient digestibility was calculated by the following equation: apparent nutrient digestibility = \[
\frac{(\text{nutrient/AIA})_{\text{diet}} - (\text{nutrient/AIA})_{\text{ digesta}}}{(\text{nutrient/AIA})_{\text{diet}}}
\] [15,25,26].

2.7. Fecal Sample Collection, DNA Extraction, 16S rRNA Gene Amplification, and Illumina HiSeq 2500 Sequencing

The fecal samples were collected on the last day of the experiment from 18 pigs, where two pigs were selected randomly within each pen (one male and one female). The feces were stored at −80 °C for subsequent analyses.

The methods in this section are similar to those used in our previous studies [27,28], in which the fecal microbial DNA was extracted using HiPure Stool DNA Kits (Magen, Guangzhou, China) according to the manufacturer’s protocols. The 16S rDNA V3–V4 region of the ribosomal RNA gene was amplified by PCR (95 °C for 2 min, followed by 27 cycles at 98 °C for 10 s, 62 °C for 30 s, and 68 °C for 30 s, with a final extension at 68 °C for 10 min) using the primers 341F: CCTACGGGNGGCWGCAG and 806R: GGCAC-TACHVGGGTATCTAAT, where the barcode is an eight-base sequence unique to each sample. PCR reactions were performed in triplicate in 50 µL mixtures containing 5 µL of 10 × KOD Buffer, 5 µL of 2.5 mM dNTPs, 1.5 µL of each primer (5 µM), 1 µL of KOD polymerase, and 100 ng of the template DNA.

Amplicons were extracted from 2% agarose gels and purified using the AxyPrep DNA Gel Extraction Kit (Axygen Biosciences, Union City, CA, USA) according to the manufacturer’s instructions and quantified using an ABI StepOnePlus Real-Time PCR System (Life Technologies, Foster City, CA, USA). Purified amplicons were pooled in an equimolar mixture and paired-end sequenced (2 × 250) on an Illumina HiSeq platform according to the standard protocols. The raw reads were deposited into the NCBI Sequence Read Archive (SRA) database (accession number: SRR9566631).

2.8. Bioinformatics Analysis

2.8.1. Read Filtering

Raw data containing adapters or low-quality reads would affect the subsequent assembly and analysis. Thus, in order to obtain high-quality clean reads, the raw reads were further filtered according to the following rules using FASTP ([https://github.com/OpenGene/fastp](https://github.com/OpenGene/fastp), accessed on 14 September 2022): (1) removing reads containing more than 10%
of unknown nucleotides (N), (2) removing reads containing less than 60% of bases with a quality (Q-value) > 20.

2.8.2. Read Assembly
Paired-end clean reads were merged as raw tags using FLSAH [29] (version 1.2.11) with a minimum overlap of 10 bp and mismatch error rates of 2%.

2.8.3. Raw Tag Filtering
Noisy sequences of raw tags were filtered by the QIIME [30] (version 1.9.1) pipeline under specific filtering conditions [31] to obtain high-quality clean tags.

2.8.4. Chimera Checking and Removal
Clean tags were searched against the reference database (http://drive5.com/uchime/uchime_download.html, accessed on 14 September 2022) to perform reference-based chimera checking using the UCHIME algorithm (http://www.drive5.com/usearch/manual/uchime_algo.html, accessed on 14 September 2022). All chimeric tags were removed, and the effective tags finally obtained were used for further analysis.

2.8.5. OTU Cluster
The effective tags were clustered into operational taxonomic units (OTUs) of ≥ 97% similarity using the UPARSE [32] pipeline. The tag sequence with the highest abundance was selected as a representative sequence within each cluster. Between-group Venn analysis was performed in R (version 3.4.1, https://cran.r-project.org/bin/windows/base/old/3.4.1/, accessed on 14 September 2022) to identify unique and common OTUs.

2.8.6. Taxonomic Classification
The representative sequences were classified into organisms by a naive Bayesian model using an RDP classifier [33] (version 2.2, http://rdp.cme.msu.edu/, accessed on 14 September 2022) based on the SILVA [34] database (https://www.arb-silva.de/, accessed on 14 September 2022), with the confidence threshold values ranging from 0.8 to 1. The abundance statistics of each taxon were visualized using Krona [35] (version 2.6, https://github.com/marbl/Krona/releases/tag/v2.6.1/, accessed on 14 September 2022). Biomarker features in each group were screened by Metastats [36] (version 20090414) and LEfSe software [37] (version 1.0, https://github.com/waldronlab/lefser, accessed on 14 September 2022).

2.8.7. Alpha Diversity Analysis
Chao1, Simpson, and all other alpha diversity indexes were calculated in QIIME. OTU rarefaction curves and rank abundance curves were plotted in QIIME. An alpha index comparison between groups was calculated by Welch’s t-test and Wilcoxon’s rank test. The alpha index comparison among the groups was computed by Tukey’s HSD test and the Kruskal–Wallis H-test.

2.9. Statistical Analysis
Data were compared among the groups using a one-way ANOVA test after normal test processing and conversion if necessary. The relative abundance of microbial communities in feces and data that did not follow a normal distribution were processed using the nonparametric Kruskal–Wallis test. Correlation analysis was performed by Pearson’s correlation tests. Significant differences were considered at \( p < 0.05 \). The initial body weight was included as a covariate in the growth performance analysis. The statistical analyses were conducted using SPSS Statistics (Version 22, https://www.ibm.com/analytics/spss-statistics-software, accessed on 14 September 2022) [27,28].
3. Results

3.1. Antibacterial Characteristics

To evaluate the antibacterial characteristics of eight TCMs, we analyzed the MICs of eight TCMs against *Escherichia coli* and *Salmonella* (Table 3). All eight TCMs had antibacterial effects on *Escherichia coli* and *Salmonella*. Compared with other TCMs, *Fructus mume* and *Scutellaria baicalensis* Georgi showed potent antibacterial properties and were then used in the subsequent preparation of fermented mixed feed.

Table 3. The MICs of the water maceration extracts of eight TCMs for *Escherichia coli* and *Salmonella*.

| TCMs          | MICs, mg mL⁻¹ |  |                  |  |
|---------------|---------------|---|------------------|---|
|               | *Escherichia coli* |  | *Salmonella*     |  |
| *Fructus mume*| 25.00         |  | 30.00            |  |
| *Scutellaria baicalensis* Georgi | 35.00         |  | 40.00            |  |
| *Rhizoma imperatae* | 130.00        |  | >250.00          |  |
| *Paconiae radix alba* | >250.00       |  | >250.00          |  |
| *Plantaginis semen* | >250.00       |  | >250.00          |  |
| *Eclipta prostrata* | >250.00       |  | >250.00          |  |
| *Fructus arctii* | >250.00       |  | >250.00          |  |
| *Portulaca oleracea* L. | >250.00       |  | >250.00          |  |

3.2. Growth Performance

The ADG, ADFI, and the F:G ratio of the experimental piglets were measured to assess their growth performance (Table 4). The ADG of the TCM and F-TCM groups increased significantly by 51.3% (*p < 0.05*) and 53.2% (*p < 0.05*) compared with the control group, respectively. Compared with the control group, the F:G ratio of the TCM and F-TCM groups decreased significantly by 33.5% (*p < 0.05*) and 26.2% (*p < 0.05*), respectively.

Table 4. The growth performance of weaned piglets (mean ± SD).

| Measure                  | Experimental Diets | p-Value |
|--------------------------|--------------------|---------|
|                          | Control             | TCM     | F-TCM   |         |
| Initial BW, kg           | 6.69 ± 0.43        | 7.96 ± 0.89 | 7.99 ± 0.93 | 0.25     |
| Final BW, kg             | 13.69 ± 0.80 b     | 18.55 ± 1.06 a | 18.71 ± 2.04 a | <0.05    |
| ADG, g                   | 259.26 ± 10.45 b   | 392.32 ± 7.16 a | 397.19 ± 9.85 a | <0.05    |
| ADFI, g                  | 535.67 ± 38.64     | 609.40 ± 14.50      | 654.44 ± 77.19    | 0.14     |
| F:G, g/g                 | 2.07 ± 0.01 a      | 1.55 ± 0.03 b      | 1.64 ± 0.08 b   | <0.05    |

* a, b, c Values within a row without a common superscript letter are significantly different (*p < 0.05*).

3.3. Diarrhea Rate

The diarrhea rate of piglets was determined according to the literature [21] (Table 5). Compared with the F-TCM group, the DM of feces decreased in control and TCM groups (*p < 0.05*). Compared with the control group, the diarrhea rate of piglets in the TCM and F-TCM groups was significantly decreased by 41.7% (*p < 0.05*) and 65.6% (*p < 0.05*), respectively.

Table 5. The DM of feces and the diarrhea rate of weaned piglets (mean ± SD).

| Measure                  | Experimental Diets | p-Value |
|--------------------------|--------------------|---------|
|                          | Control             | TCM     | F-TCM   |         |
| Feces, DM %              | 60.16 ± 4.89 b     | 63.15 ± 3.80 b | 67.34 ± 2.95 a | <0.05    |
| Diarrhea rate, %         | 22.22 ± 4.13 a     | 12.96 ± 3.23 b | 7.66 ± 2.26 c | <0.05    |

* a, b, c Values within a row without a common superscript letter are significantly different (*p < 0.05*).
3.4. Apparent Nutrient Digestibility

To determine whether TCM supplementation could improve the apparent nutrient digestibility in weaned piglets, we assessed the digestibility of DM, CP, EE, CF, ash, and NFE (Table 6). The apparent digestibility of DM and EE increased in the TCM and F-TCM groups in comparison with the control group \((p < 0.05)\), while no significant difference was noted between the TCM and F-TCM groups. There was no significant difference in the apparent digestibility of CP, CF, ash, or NFE among all the groups \((p > 0.05)\).

Table 6. The apparent nutrient digestibility of weaned piglets (mean ± SD).

| Nutrient | Control | TCM | F-TCM | \(p\)-Value |
|----------|---------|-----|-------|------------|
| DM, %    | 62.93 ± 5.12 \(^b\) | 70.20 ± 0.99 \(^a\) | 66.59 ± 4.66 \(^a\) | <0.05 |
| CP, %    | 64.48 ± 5.67 | 68.86 ± 4.16 | 72.93 ± 6.63 | 0.26 |
| EE, %    | 79.05 ± 3.75 \(^b\) | 89.93 ± 6.48 \(^a\) | 85.09 ± 4.34 \(^a\) | <0.05 |
| CF, %    | 72.14 ± 4.23 | 86.57 ± 5.34 | 62.77 ± 2.65 | 0.18 |
| Ash, %   | 85.82 ± 4.71 | 86.67 ± 3.34 | 87.21 ± 5.72 | 0.94 |
| NFE, %   | 85.53 ± 5.46 | 68.25 ± 3.81 | 74.93 ± 3.37 | 0.54 |

\(^a,b\) Values within a row without a common superscript letter are significantly different \((p < 0.05)\).

3.5. The Composition of Fecal Microbiota

At a cutoff level of 3%, no effect on the ACE and Chao richness estimators was observed in all groups (Figure 1). The Shannon diversity estimator in the F-TCM group was significantly increased \((p < 0.05)\), while the Simpson index decreased compared with the control and TCM groups \((p < 0.01)\).

The OTU distribution of the microbial communities of the different treatment groups had a certain degree of similarity and specificity. In order to understand the species differences, Venn diagrams were used to show the common and unique information among the different groups based on the OTU abundance information of the samples. As shown in Figure 2, 549 microbial species were common to all groups; however, 101 OTUs were unique in the control, and 136 and 146 OTUs were unique in the TCM and F-TCM groups, respectively.

At the phylum level, Bacteroidetes and Firmicutes were the predominant phyla in the fecal microbiota of piglets, with a total abundance of >95%, followed by the phyla Actinobacteria and Proteobacteria (Figure 3A,D). The abundance statistics of each taxon were visualized using Krona, the total profiling of the composition of microbial species is shown in Figure S1, and the composition of microbial species at the phylum level of Firmicutes (Figure S2), Bacteroidetes (Figure S3), and Actinobacteria (Figure S4) are presented in supplementary materials.

At the genus level, in the phylum Bacteroidetes, 10 genera with a relative abundance of >1% were found to be dominant (Figure 3E). In the phylum Firmicutes, 17 genera with a relative abundance of >1% were found to be dominant (Figure 3F). Among these, compared with the control group, the relative abundance of Erysipelotrichaceae_UCG-004 and Ruminococcaceae_UCG-004 were increased, and the abundance of Prevotellaceae_UCG-003 and Oscillospira were decreased in the TCM and F-TCM groups \((p < 0.05)\), Figure 4. Compared with the control and F-TCM groups, the abundance of Acidaminococcus, Prevotella, and Megaplasma were increased, while the abundance of Bifidobacterium, Coprococcus, Lachnospiraceae, and Ruminococcaceae were decreased in the TCM group \((p < 0.05)\), Figure 4. Compared with the control and TCM groups, the relative abundance levels of Coprococcus, Lachnospiraceae, and Ruminococcaceae were decreased in the F-TCM group \((p < 0.05)\), Figure 4.
the *Prevotellaceae*_NK3B31 group, *Faecalibacterium*, *Oscillibacter*, and *Ruminococcaceae_UCG-008* were increased, while the abundance of *Acidaminococcus*, *Bifidobacterium*, *Prevotella_7*, and *Megasphaera* were decreased in the F-TCM group (*p* < 0.05, Figure 4).

At the OTU level, compared with the control group, the relative abundance of *Ruminococcaceae_UCG-014*-related OTUs (OTU157 and OTU043), the *Subdoligranulum*-related OTU214, and the *Prevotellaceae*-related OTU292 were increased, while the abundance of the *Ruminococcaceae_UCG-002*-related OTU037 and the *Prevotella_9*-related OTU295 were decreased in the TCM and F-TCM groups (*p* < 0.05, Figure 5). Compared with the control and F-TCM groups, the abundance of the *Christensenellaceae_R-7* group-related OTU210, the *Acidaminococcus*-related OTU112, and the *Prevotella_7*-related OTU007 were increased, while the abundance of the *Candidatus Soleaferrea*-related OTU133, the *Oscillibacter*-related OTU300, the *Oscillospira*-related OTUs (OTU242 and OTU041), the *Ruminiclostridium_9*-related OTU202, the *Prevotella_1*-related OTU069, the *Prevotella_9*-related OTUs (OTU131 and OTU057), and the *Prevotellaceae_UCG-003*-related OTU012 were decreased in the TCM group (*p* < 0.05, Figure 5). Compared with the TCM group, the abundance of the *Coprococcus_1*-related OTU315, the *Lachnoclostridium*-related OTU148, the *Candidatus Soleaferrea*-related OTU133, the *Oscillibacter*-related OTU300, the *Oscillospira*-related OTUs (OTU242 and OTU041), the *Ruminiclostridium_9*-related OTU202, the *Ruminococcaceae_UCG-005*-related OTU180, the *Ruminococcaceae_UCG-014*-related OTUs (OTU354, OTU184, OTU157, and OTU043), the *Prevotella_1*-related OTU069, and the *Prevotella_9*-related OTUs (OTU131 and OTU057) were increased, while the abundance of the *Ruminococcaceae_UCG-002*-related OTU037, the *Acidaminococcus*-related OTU112, and the *Prevotella_7*-related OTU007 were decreased in the F-TCM group (*p* < 0.05, Figure 5).
Figure 2. Venn diagram analysis of the OTUs among the treatment groups.

Figure 3. The composition and structure of the fecal microbiota in weaned piglets (relative abundance of more than 1%). The Bacteroidetes and Firmicutes phyla constituted approximately 95% of the identified sequences (A), followed by Actinobacteria and Proteobacteria as follows: five families (B) and 10 genera (E) in Bacteroidetes; eight families (C) and 17 genera (F) in Firmicutes; one family (D) in Actinobacteria; two families (D) in Proteobacteria. P: phylum.
Figure 4. The relative abundance of the fecal microbiota of weaned piglets at the genus level. F: family. * indicate significant differences among three groups ($p < 0.05$), ** indicate significant differences among three groups ($p < 0.01$).
Figure 5. The relative abundance of the fecal microbiota of weaned piglets at the OTU level. F: family; G: genus. * indicate significant differences among three groups ($p < 0.05$), ** indicate significant differences among three groups ($p < 0.01$).

3.6. Correlation between Fecal Microbiota and Apparent Nutrient Digestibility

In order to investigate the relationship between the intestinal microbial community and apparent nutrient digestibility in piglets, the correlations were analyzed (Table 7 and Figure 6).

Table 7. Correlations of fecal microbial richness and diversity estimators with apparent nutrient digestibility.

| Apparent Digestibility, % | Correlation Coefficient |
|---------------------------|-------------------------|
|                           | ACE         | Chao       | Shannon    | Simpson    |
| DM                       | 0.03 *      | 0.05 *     | 0.53       | 0.16       |
| CP                       | 0.34        | 0.28       | 0.82       | 0.97       |
| EE                       | 0.89        | 0.71       | 0.03 *     | 0.02 *     |
| CF                       | 0.67        | 0.82       | 0.10       | 0.17       |
| Ash                      | 0.74        | 0.61       | 0.82       | 0.59       |
| NFE                      | 0.34        | 0.72       | 0.44       | 0.44       |

* Significant correlation ($p < 0.05$).

As shown in Table 7, the microbial richness estimators (ACE, Chao) were positively correlated with apparent DM digestibility ($p < 0.05$), and the microbial diversity indices (Shannon, Simpson) were positively correlated with apparent EE digestibility ($p < 0.05$).
Correlation analysis of fecal microbiota (genus and OTU levels) and apparent nutrient digestibility: (A) genus level; (B) OTU level. The color indicates the Pearson coefficient distribution: red represents a positive correlation \((p < 0.05)\), blue represents a negative correlation \((p < 0.05)\), and white shows that the correlation was not significant \((p > 0.05)\). F: family; G: genus.

At the genus level (Figure 6A), the relative abundance of the *Ruminococcaceae* _UCG-014_ and *Lachnospiraceae* _FCS020_ groups were negatively correlated with apparent DM digestibility \((p < 0.05)\); the abundance of *Ruminococcaceae* _UCG-014_ and *Lachnolostrium* was negatively correlated with apparent EE digestibility \((p < 0.05)\); and the abundance of *Oscillospora* was negatively correlated with apparent ash digestibility \((p < 0.05)\).

At the OTU level (Figure 6B), the relative abundance of the *Ruminococcaceae* _UCG-014_-related OTUs (OTU043 and OTU157) and the family *Prevotellaceae*-related OTU371 were positively correlated, while the abundance of the family *Ruminococcaceae*-related OTU037 and the *Prevotella*_9-related OTU295 was negatively correlated with the apparent DM digestibility \((p < 0.05)\). The abundance of the *Subdoligranulum*-related OTU214 was negatively correlated with the apparent CP digestibility \((p < 0.05)\); the abundance of the *Prevotella*_9-related OTU295, the family *Ruminococcaceae*-related OTUs (OTU103 and OTU165), and the family *Bacteroidale_S24-7* group-related OTU113 was positively correlated with apparent EE digestibility, while the abundance of the *Ruminococcaceae* _UCG-014_-related OTUs (OTU157, OTU354, and OTU180), the *Prevotellaceae*_NK3B31 group-related OTUs (OTU056, OTU065, OTU116, and OTU188), the *Prevotella*_1-related OTU069, the family *Ruminococcaceae*-related OTU292, and the *Lachnolostrium*-related OTU148 was negatively correlated with the apparent EE digestibility \((p < 0.05)\). The abundance of *Oscillospora*-related OTU041, *Prevotellaceae*_NK3B31 group-related OTU188, and family *Prevotellaceae*-related OTU292 were positively correlated with apparent CF digestibility \((p < 0.05)\).

### 4. Discussion

With the implementation of the policy to remove AGPs in animal production in China, natural plants and their application as AGP substitutes have gained increasing interest in the research community because of their safety, efficiency, and availability [38]. TCMs are considered better alternatives for improving animal health and resisting infectious diseases. *Fructus mume* (“wumei” in Chinese) has long been used in China to treat chronic coughs, expectoration, ulceration, chronic diarrhea, and gastrointestinal diseases [39–44]. This medicinal effect is due to its antioxidant [44], antibacterial [45], and anti-inflammatory properties [43,44], and its protective ability against gastrointestinal diseases via the opsonization of intestinal commensal bacteria, as well as its ability to alleviate epithelial injury and inflammation [39]. *Scutellaria baicalensis* Georgi (“huangqin” in Chinese) is also an old
and well-known component of TCM and is widely used for the treatment of bronchitis, hepatitis, tumors, and inflammatory diseases [46–50]. Numerous research studies have indicated that the therapeutic effects of *Scutellaria baicalensis* Georgi are due to its various pharmacological activities, including its antiangiogenic, anti-inflammatory, antimicrobial, immunoenhancing, and antioxidative effects [51–54]. Very little was found in the literature on the effects of dietary TCM in weaned piglets. Our study systematically investigated the data and aimed to ascertain the effects of these two TCM feed additives on the growth performance, apparent nutrient digestibility, and fecal microbiota of weaned piglets.

Prior studies have reported that fermented feed significantly increases the body weight and ADG of piglets [55], laying hen chicks [56], and geese [57]. Several reports have shown that TCM additives significantly improved the final BW, ADG, and FCR in lambs and hogs [12] and promoted growth performance in heat-stressed beef cattle, which was associated with better physiological status [11]. These findings are contrary to a previous study showing that dietary TCM led to greater feed intake but no significant differences in the final BW, ADG, or F:G ratio [58]. There were few reports on the effects of fermentation with TCM mixtures on the growth performance of weaned piglets. In our study, dietary supplementation with *Fructus mume* and *Scutellaria baicalensis* Georgi, fermented or not fermented, led to no significant differences in ADFI during the experiment period. The findings of the current study do not support previous research where the authors suggested that the inclusion of supplemental TCMs may improve pigs’ appetite [58]. The results of this study showed that the final BW and ADG increased, while the F:G ratio and diarrhea rate decreased in weaned piglets in the TCM and F-TCM groups, suggesting that *Fructus mume* and *Scutellaria baicalensis* Georgi supplementation in the diet improves growth performance, leading to greater weight gain and improved health [10–12]. However, fermentation with *Fructus mume* and *Scutellaria baicalensis* Georgi in the diet had no significant effect on the growth performance of piglets.

One interesting finding in our study was observed in the TCM and F-TCM groups, in which ADG was increased while ADFI was not significantly changed; considerably more work will need to be carried out to determine apparent nutrient digestibility in piglets. Previous studies have explored whether dietary TCM increases the apparent digestibility of CP in finishing pigs [39] and CP, Ca, P, and NDF in weaned piglets [60]. Dietary TCM has also been suggested to improve the apparent digestibility of OM, CP, ADF, Ca, and P in beef cattle, even under heat stress [10,11]. However, the findings of the current study do not support the previous research; as shown in Table 6, there was no significant difference in the apparent digestibility of CP or CF among the three groups. In addition, the partial substitution of fermented feed in the diet increased the apparent digestibility of CP and CF in growing-finishing pigs [61]; CP and EE in Xuefeng black-boned chickens [62]; and DM, CP, CF, NDF, and ADF in lactating dairy cows [63]. Data on fermented TCM feed are lacking; in our study, feed fermented with TCM increased the apparent digestibility of DM and EE compared with the non-fermented group (TCM group), suggesting that fermented feed with *Fructus mume* and *Scutellaria baicalensis* Georgi mixture increased the digestibility of DM and EE in weaned piglets.

Recently, studies have found that the gut microbiota may explain the therapeutic effects of TCM [6,64]. An increasing number of studies have investigated the interactions between the gut microbiota and TCM, suggesting that the gut microbiota can directly affect the absorption, metabolism, and pharmacological activity of TCM [2,3,65]. In our study, the composition of fecal microbiota in weaned piglets was analyzed. Dietary *Fructus mume* and *Scutellaria baicalensis* Georgi decreased the diversity but did not have a significant effect on the richness of fecal microbiota; this finding is inconsistent with that of Zou (2021), who reported that a Huangqin decoction (HQD) could increase the diversity of the intestinal microbiota of cholestatic mice [66]. This inconsistency may be due to the different species. The current study also found that feed fermented with these two TCMs improved the diversity of fecal microbiota in piglets. Previous results suggested that eight Chinese herbs (Chinese name: “jian ji san”) fermented with *Zygosaccharomyces rouxii* and
their fermentation products increased the diversity of the foregut microbial community of broiler chickens [67], which is in agreement with our results. According to these data, we can infer that dietary *Fructus mume* and *Scutellaria baicalensis* Georgi affected the fecal microbial composition by altering its diversity in weaned piglets.

In our study, *Bacteroidetes* and *Firmicutes* were the predominant phyla in the fecal microbiota of piglets, similar to the findings of previous studies [28,68,69]. As mentioned in the literature review, the gut microbial mediation of the potential therapeutic mechanism of TCMs can be attributed to the production of short-chain fatty acids (SCFAs) [67,70], mostly acetic, propionic, and butyric acids, which play an important role in maintaining the intestinal health of pigs [70,71]. In Figure 4, there is a clear trend of the increased relative abundance of *Acidaminococcus*, *Prevotella*, and *Megasphaera* in the TCM group and a decreased abundance in the F-TCM group; all three genera are considered to be SCFA-producing bacteria [70,72], suggesting that the diet supplemented with *Fructus mume* and *Scutellaria baicalensis* Georgi increased the abundance of SCFA-producing bacteria in the hindgut, which may have promoted the intestinal health of the weaned piglets. However, the prefermentation of these two TCMs, dietary fibers, and other indigestible carbohydrates led to their degradation before being fed to the piglets [70], explaining the decreasing tendency of apparent CF digestibility and the abundance of *Acidaminococcus*, *Prevotella*, and *Megasphaera* in the F-TCM group. Another important finding in the current study was that the relative abundance of *Coprococcus*, *Lachnospiraceae_FCS020 group*, and *Oscillibacter* were increased in the F-TCM group; these genera have been identified as the most active and healthy microbiome constituents in the intestinal environment in healthy adult humans and animals [28,73–75]. Thus, these findings suggest that fermentation with *Fructus mume* and *Scutellaria baicalensis* Georgi results in the improved healthy intestinal flora, which, in turn, could favor the intestinal health of weaned piglets.

Another important objective of this study was to investigate the relationship between the gut microbial characteristics and the apparent nutrient digestibility in weaned piglets, so the correlation between the apparent nutrient digestibility and significant microbial genera and OTUs were further analyzed. Significant correlations between microbial diversity and apparent EE digestibility were observed, suggesting a potential link between changes in the intestinal flora and apparent EE digestibility. At the OTU level, the relative abundance of 10 OTUs that increased in the F-TCM group was positively correlated with apparent EE digestibility, while four OTUs for which the abundance was decreased in the F-TCM group were negatively correlated with apparent EE digestibility. Within these OTUs, most belong to the families *Ruminococcaceae* and *Prevotellaceae*, which are considered to be the core bacteria detected in 99% of fecal samples obtained from commercial swine worldwide [76]. Prior studies have shown that the abundance of *Ruminococcaceae* is negatively correlated with high-fat-diet (HFD)-induced obesity in a ripened pu-erh tea extract (PETe) intervention in mice [77], while the relative abundance of *Prevotellaceae* and *Prevotellaceae_NK3B31_group* was observed to increase through supplementation with pea seed coats (PSCs) and stachyose in mice and rats fed an HFD [78,79]. According to these findings, we can infer that the digestion and absorption of nutritional lipids in the diet were closely related to the changes in the *Ruminococcaceae* and *Prevotellaceae* families. Our correlation results are consistent with previous studies focusing on the gut microbiota and the apparent nutrient digestibility of grower pigs and sows [15,80] and support the possible relationship between the gut microbiota and the regulation of dietary nutrient utilization in piglets.

5. Conclusions

Dietary supplementation with *Fructus mume* and *Scutellaria baicalensis* Georgi in antibiotic-free feed improved the final BW and ADG, increased the abundance of SCFA-producing bacteria in the hindgut, and decreased the F:G ratio and diarrhea rate, yielding healthier weaned piglets that gained more weight. Fermentation with these TCMs enhanced the apparent digestibility of DM and EE and improved the healthy intestinal flora,
which, in turn, could favor the intestinal health of weaned piglets. There was a significant correlation between the increased apparent EE digestibility in the TCM diets and the diversity of fecal microbiota. Dietary TCMs affect the fecal microbial composition by changing the abundance of certain genera belonging to the Ruminococcaceae, Prevotellaceae, and Lachnospiraceae families, which may further increase the apparent EE digestibility of weaned piglets. Nevertheless, our study cannot demonstrate causality, and further experimental studies are needed to address this.

**Supplementary Materials:** The following supporting information can be downloaded at: https://www.mdpi.com/article/10.3390/ani12182418/s1, The supplementary result for the fermented feed physiological characteristics is shown in Table S1. The supplementary results for Krona analysis are shown in Figures S1–S4. Table S1: The pH values at different fermentation times; Figure S1: The total profiling of the composition of microbial species; Figure S2: The composition of microbial species at the phylum level of Firmicutes; Figure S3: The composition of microbial species at the phylum level of Bacteroidetes; Figure S4: The composition of microbial species at the phylum level of Actinobacteria.

**Author Contributions:** Conceptualization, F.Z., E.J., X.L., X.J. and H.H.; methodology, F.Z., E.J., X.J. and H.H.; formal analysis, F.Z. and X.J.; resources, F.Z.; data curation, F.Z., X.J. and H.H.; writing—original draft preparation, F.Z.; writing—review and editing, F.Z., X.L. and E.J.; project administration, F.Z. and X.J.; funding acquisition, F.Z. and E.J. All authors have read and agreed to the published version of the manuscript.

**Funding:** This work was supported by the National Natural Science Foundation of China (32172816), the Anhui Natural Science Foundation Project (2108085MC117, 2208085MC77), the Natural Science Key Foundation of Anhui Education Department (KJ2021A0868), the Laboratory Open Project of Anhui Province Key Laboratory of Animal Nutrition Regulation and Health (APKLNRH202001), the High-level Talents Introduction Foundation of Anhui Science and Technology University (DKYJ202101), the College Student Innovation and Entrepreneurship Project (202110879058, S202110879172), and the Special Fund for Anhui Agriculture Research System (AHCYJSTX-05-16).

**Institutional Review Board Statement:** This study was conducted according to the guidelines for the care and use of experimental animals of the Ministry of Science and Technology of the People’s Republic of China (Approval Number: 2006-398) and approved by the Experimental Animal Ethical Committee of Anhui Science and Technology University. The piglets came from a local family farm, and the farm owner was informed and agreed to the implementation of this experimental protocol.

**Informed Consent Statement:** The farm owner was informed and agrees to the implementation of this experimental protocol, including the composition of feed and sample collection for piglets.

**Data Availability Statement:** Raw data collected and presented in this study are available on request from the corresponding author.

**Acknowledgments:** The authors would like to thank You Liu and Yanfen Liu from Guangdong Ocean University for her assistance with the administration of the experiment.

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

**References**

1. Feng, W.; Liu, J.; Huang, L.; Tan, Y.; Peng, C. Gut microbiota as a target to limit toxic effects of traditional Chinese medicine: Implications for therapy. *Biomed. Pharm. 2021, 133, 111047. [CrossRef] [PubMed]*

2. Zheng, Y.; Ding, Q.; Wei, Y.; Gou, X.; Tian, J.; Li, M.; Tong, X. Effect of traditional Chinese medicine on gut microbiota in adults with type 2 diabetes: A systematic review and meta-analysis. *Phytomedicine 2021, 88, 153455. [CrossRef] [PubMed]*

3. Gong, X.; Li, X.; Bo, A.; Shi, R.Y.; Li, Q.Y.; Lei, L.J.; Zhang, L.; Li, M.H. The interactions between gut microbiota and bioactive ingredients of traditional Chinese medicines: A review. *Pharmacol. Res. 2020, 157, 104824. [CrossRef]*

4. Zhang, X.; Yang, Y.; Zhang, F.; Yu, J.; Sun, W.; Wang, R.; Wu, C. Traditional Chinese medicines differentially modulate the gut microbiota based on their nature (Yao-Xing). *Phytomedicine 2021, 85, 153496. [CrossRef] [PubMed]*

5. Zhang, G.X.; Jin, L.; Jin, H.; Zheng, G.S. Influence of Dietary Components and Traditional Chinese Medicine on Hypertension: A Potential Role for Gut Microbiota. *Evid.-Based Complement. Alternat. Med. 2021, 2021, 5563073. [CrossRef] [PubMed]*

6. Lu, Y.; Liu, H.; Yang, K.; Mao, Y.; Meng, L.; Yang, L.; Ouyang, G.; Liu, W. A comprehensive update: Gastrointestinal microflora, gastric cancer and gastric premalignant condition, and intervention by traditional Chinese medicine. *J. Zhejiang Univ. Sci. B 2022, 23, 1–18. [CrossRef]*
Animals 2022, 12, 2418

7. Liu, Y.; Li, B.G.; Su, Y.H.; Zhao, R.X.; Song, P.; Li, H.; Cui, X.H.; Gao, H.M.; Zhai, R.X.; Fu, X.J.; et al. Potential activity of Traditional Chinese Medicine against Ulcerative colitis: A review. J. Ethnopharmacol. 2022, 289, 115084. [CrossRef]

8. Zhao, H.; He, M.; Zhang, M.; Sun, Q.; Zeng, S.; Chen, L.; Yang, H.; Liu, M.; Ren, S.; Meng, X.; et al. Colorectal Cancer, Gut Microbiota and Traditional Chinese Medicine: A Systematic Review. Am. J. Chin. Med. 2021, 49, 805–828. [CrossRef]

9. Feng, W.; Ao, H.; Peng, C.; Yan, D. Gut microbiota, a new frontier to understand traditional Chinese medicines. Pharmacol. Res. 2019, 142, 176–191. [CrossRef]

10. Song, X.; Luo, J.; Fu, D.; Zhao, X.; Bunlue, K.; Xu, Z.; Qu, M. Traditional chinese medicine prescriptions enhance growth performance of heat stressed beef cattle by relieving heat stress responses and increasing apparent nutrient digestibility. Asian-Australas. J. Anim. Sci. 2014, 27, 1513–1520. [CrossRef]

11. Chen, J.; Guo, K.; Song, X.; Lan, L.; Liu, S.; Hu, R.; Luo, J. The anti-heat stress effects of Chinese herbal medicine prescriptions and rumen-protected gamma-aminobutyric acid on growth performance, apparent nutrient digestibility, and health status in beef cattle. Anim. Sci. J. 2020, 91, e13361. [CrossRef] [PubMed]

12. Du, Z.; Risu, N.; Gentu, G.; Jia, Y.; Cai, Y. Growth performance, apparent digestibility, and N balance in Mongolian lambs and hogs fed diets supplemented with a Chinese traditional herbal medicine complex. Anim. Sci. J. 2018, 89, 1451–1458. [CrossRef] [PubMed]

13. Sun, Z.; Li, H.; Li, Y.; Qiao, J. Lactobacillus salivarius, a potential probiotic to improve the health of LPS-challenged piglet intestine by alleviating inflammation as well as oxidative stress in a dose-dependent manner during weaning transition. Front. Vet. Sci. 2020, 7, 547425. [CrossRef]

14. Dyar, O.J.; Zhang, T.; Peng, Y.; Sun, M.; Sun, C.; Yin, J.; Ding, L.; Sun, C.; Wang, Y.; Sun, Q.; et al. Knowledge, attitudes and practices relating to antibiotic use and antibiotic resistance among backyard pig farmers in rural Shandong province, China. Prev. Vet. Med. 2020, 175, 104858. [CrossRef]

15. Niu, Q.; Li, P.; Hao, S.; Kim, S.W.; Du, T.; Hua, J.; Huang, R. Characteristics of gut microbiota in sows and their relationship with apparent nutrient digestibility. Int. J. Mol. Sci. 2019, 20, 870. [CrossRef]

16. Agga, G.E.; Scott, H.M.; Amachawadi, R.G.; Nagaraja, T.; Vinasco, J.; Bai, J.; Norby, B.; Renter, D.G.; Dritz, S.S.; Nelssen, J.L.; et al. Effects of chlortetracycline and copper supplementation on antimicrobial resistance of fecal Escherichia coli from weaned pigs. Prev. Vet. Med. 2014, 114, 231–246. [CrossRef]

17. Zou, X.; Weng, M.; Ji, X.; Guo, R.; Zheng, W.; Yao, W. Comparison of antibiotic resistance and copper tolerance of Enterococcus spp. and Lactobacillus spp. isolated from piglets before and after weaning. J. Microbiol. 2017, 55, 703–710. [CrossRef]

18. CLSI. Methods for Dilution Antimicrobial Susceptibility Tests for Bacteria That Grow Aerobically: Approved Standard-Ninth Edition; Wayne, P., Ed.; Clinical and Laboratory Standards Institute: Wayne, PA, USA, 2012; p. CLSI document M07-A9.

19. NRC. Nutrient Requirements of Swine, 11th ed.; Whitacre, P.T., Ed.; The National Academies Press: Washington, DC, USA, 2012.

20. Yu, J.; Song, Y.; Yu, B.; He, J.; Zheng, P.; Mao, X.; Huang, Z.; Luo, Y.; Luo, J.; Yan, H.; et al. Tannic acid prevents post-weaning diarrhea by improving intestinal barrier integrity and function in weaned pigs. J. Anim. Sci. Biotechnol. 2020, 11, 87. [CrossRef]

21. Yin, L.; Li, J.; Wang, H.; Yi, Z.; Wang, L.; Zhang, S.; Li, X.; Wang, Q.; Li, J.; Yang, H.; et al. Effects of vitamin B6 on the growth performance, intestinal morphology, and gene expression in weaned piglets that are fed a low-protein diet. J. Anim. Sci. 2020, 98, skaa022. [CrossRef]

22. Fouhse, J.M.; Gao, J.; Vasanthan, T.; Izydorczyk, M.; Beattie, A.D.; Zijlstra, R.T. Whole-grain fiber composition influences site of nutrient digestion, standardized ileal digestibility of amino acids, and whole-body energy utilization in grower pigs. J. Nutr. 2017, 147, 29–36. [CrossRef]

23. Ngoc, T.T.; Len, N.T.; Lindberg, J.E. Impact of fibre intake and fibre source on digestibility, gut development, retention time and growth performance of indigenous and exotic pigs. Animal 2013, 7, 736–745. [CrossRef] [PubMed]

24. Xie, Y.; Zhang, Q.; Wang, L.; Wang, Y.; Cheng, Z.; Yang, Z.; Yang, W. The effects of partially or completely substituted dietary zinc sulfate by lower levels of zinc methionine on growth performance, apparent total tract digestibility, immune function, and visceral indices in weaned pigs. Animals 2019, 9, 236. [CrossRef] [PubMed]

25. Latimer, G.W. Official Methods of Analysis of AOAC International, 21st ed.; Latimer, G.W., Ed.; AOAC: Rockville, MD, USA, 2019.

26. Yu, M.; Zhang, C.; Yang, Y.; Mu, C.; Su, Y.; Yu, K.; Zhu, W. Long-term effects of early antibiotic intervention on blood parameters, apparent nutrient digestibility, and fecal microbial fermentation profile in pigs with different dietary protein levels. J. Anim. Sci. Biotechnol. 2017, 8, 60–72. [CrossRef] [PubMed]

27. Zhang, F.; Zheng, W.; Guo, R.; Yao, W. Effect of dietary copper level on the gut microbiota and its correlation with serum inflammatory cytokines in Sprague-Dawley rats. J. Microbiol. 2017, 55, 694–702. [CrossRef] [PubMed]

28. Zhang, F.; Zheng, W.; Xue, Y.; Yao, W. Suhuai sucking piglet hindgut microbiome-metabolome responses to different dietary copper levels. Appl. Microbiol. Biotechnol. 2019, 103, 853–868. [CrossRef]

29. Magoc, T.; Salzberg, S.L. FLASH: Fast length adjustment of short reads to improve genome assemblies. Bioinformatics 2011, 27, 2957–2963. [CrossRef]

30. Caporaso, J.G.; Kuczynski, J.; Stombaugh, J.; Bittinger, K.; Bushman, F.D.; Costello, E.K.; Fierer, N.; Pena, A.G.; Goodrich, J.K.; Gordon, J.I.; et al. QIIME allows analysis of high-throughput community sequencing data. Nat. Methods 2010, 7, 335–336. [CrossRef]

31. Bokulich, N.A.; Subramanian, S.; Faith, J.J.; Gevers, D.; Gordon, J.I.; Knight, R.; Mills, D.A.; Caporaso, J.G. Quality-filtering vastly improves diversity estimates from Illumina amplicon sequencing. Nat. Methods 2013, 10, 57–59. [CrossRef]
Animals 2022, 12, 2418

18 of 19

32. Edgar, R.C. UPARSE: Highly accurate OTU sequences from microbial amplicon reads. Nat. Methods 2013, 10, 996–998.

33. Wang, Q.; Garrity, G.M.; Tiedje, J.M.; Cole, J.R. Naive Bayesian classifier for rapid assignment of rRNA sequences into the new bacterial taxonomy. Appl. Environ. Microbiol. 2007, 73, 5261–5267. [CrossRef]

34. Puersse, E.; Quast, C.; Knittel, K.; Fuchs, B.M.; Ludwig, W.; Fleisch, J.; Gockner, F.O. SILVA: A comprehensive online resource for quality checked and aligned ribosomal RNA sequence data compatible with ARB. Nucleic Acids Res. 2007, 35, 7188–7196. [CrossRef][PubMed]

35. Ondov, B.D.; Bergman, N.H.; Phillippy, A.M. Interactive metagenomic visualization in a Web browser. BMC Bioinform. 2011, 12, 385–394. [CrossRef][PubMed]

36. White, J.R.; Nagarajan, N.; Pop, M. Statistical methods for detecting differentially abundant features in clinical metagenomic samples. PloS Comput. Biol. 2009, 5, e1000352. [CrossRef][PubMed]

37. Segata, N.; Izard, J.; Waldron, L.; Gevers, D.; Miropolsky, L.; Garrett, W.S.; Huttenhower, C. Metagenomic biomarker discovery and explanation. Genome Biol. 2011, 12, R60. [CrossRef]

38. Jin, B.R.; Chung, K.S.; Kim, H.J.; An, H.J. Chinese Skullcap (Scutellaria baicalensis Georgi) inhibits inflammation and proliferation of rat gastric epithelial cells induced by H. pylori. J. Ethnopharmacol. 2007, 108, 61–67. [CrossRef][PubMed]

39. Melas, J.; Ntini, E.; Davalos, M.; Jimenez, E.; Rosado, B.; Marques, J.; Ferreira, C.; Gomes, C.; Calzada, D.; Polanczyk, C.; et al. Protective effects of Fructus mume extracts on osteoporosis. PLoS One 2017, 12, e0169819. [CrossRef][PubMed]

40. Xu, Z.; Wang, Q.; Zhang, X.; Wang, W.; Zhang, D.; Ma, Y.; Zhang, D.; Chen, M. Fructus Mume (Wu Mei) Attenuates Acetic Acid-Induced Ulcerative Colitis by Regulating Inflammatory Cytokine, Reactive Oxygen Species, and Neuropeptide Levels in Model Rats. J. Med. Food 2022, 25, 389–401. [CrossRef]

41. Xu, Z.; Zhang, X.; Lu, R.; Zhang, D.; Zou, T.; Chen, M.; Zhang, D. Mechanism of Fructus Mume Pills Underlying Their Protective Effects in Rats with Acetic Acid-Induced ulcerative Colitis via the Regulation of Inflammatory Cytokines and the VEGF-PI3K/Akt-eNOS Signaling Pathway. Evid.-Based Complement. Alternat. Med. 2022, 2022, 462131. [CrossRef]

42. Liu, Z.; Feng, Y.; Ma, P.; Fan, L.; Zhao, L.; Wang, M.; Li, X. An integrated strategy for anti-inflammatory quality markers screening of traditional Chinese herbal medicine Mume Fructus based on psychochemical analysis and anti-colitis activity. Phytother. Res. 2022, 39, 99–119. [CrossRef]

43. Xiang, J.; Liu, X.; Zhong, S.; Fang, Z.; Shen, S.; Tang, J.; Lai, S.; Lai, K. Fructus mume protects against cigarette smoke induced chronic cough guinea pig. J. Med. Food 2020, 23, 191–197. [CrossRef]

44. Kim, M.S.; Bang, J.H.; Lee, J.; Han, J.S.; Kang, H.W.; Jeon, W.K. Fructus mume ethanol extract prevents inflammation and normalizes the seNhippocampal cholinergic system in a rat model of chronic cerebral hypoperfusion. J. Med. Food 2016, 19, 196–204. [CrossRef]

45. Xing, H.; Zhang, L.; Ma, J.; Liu, Z.; Song, C.; Liu, Y. Fructus mume extracts alleviate diarrhea in breast cancer patients receiving the combination therapy of lapatinib and revlimid. Front. Pharmacol. 2019, 8, 516. [CrossRef][PubMed]

46. Jeon, W.K.; Ma, J.; Choi, B.R.; Han, S.H.; Jin, Q.; Hwang, B.Y.; Han, J.S. Effects of Fructus mume extract on MAPK and NF-kappaB signaling and the resultant improvement in the cognitive deficits induced by chronic cerebral hypoperfusion. Evid.-Based Complement. Alternat. Med. 2012, 2012, 408538. [CrossRef][PubMed]

47. Feng, T.; Zhou, L.; Gai, S.; Zhai, Y.; Gou, N.; Wang, X.; Zhang, X.; Cui, M.; Wang, L.; Wang, S. Acacia catechu (L.) Wild and Scutellaria baikalensis Georgi extracts suppress LPS-induced pro-inflammatory responses through NF-small k, CrysicilB, MAPK, and PI3k-Akt signaling pathways in alveolar epithelial type II cells. Phytother. Res. 2019, 33, 3251–3260. [CrossRef][PubMed]

48. Chen, P.; Zhou, X.; Zhang, L.; Shan, M.; Bao, B.; Cao, Y.; Kang, A.; Ding, A. Anti-inflammatory effects of Huangqin tang extract in mice on ulcerative colitis. J. Ethnopharmacol. 2015, 162, 207–214. [CrossRef]

49. Zhao, F.; Chang, Y.; Gao, L.; Qin, X.; Du, G.; Zhang, X.; Zhou, Y. Protective effects of Scutellaria baicalensis Georgi extract on D-galactose induced aging rats. Metab. Brain Dis. 2018, 33, 1401–1412. [CrossRef]

50. Zhou, J.; Luo, Y.; Kang, X.; Bian, F.; Liu, D. The root extract of Scutellaria baikalensis Georgi promotes beta cell function and protects from apoptosis by inducing autophagy. J. Ethnopharmacol. 2022, 284, 114790. [CrossRef]

51. Wang, Y.; Cai, Y.; Li, F.; Zhang, M.; Wu, Y.; Dai, Y.; Zheng, F.; Yue, H.; Bai, B. Effects of Scutellaria baikalensis Georgi on intestinal flora in rats with spleen deficiency and damp-heat. J. Pharm. Biomed. Anal. 2022, 217, 114831. [CrossRef][PubMed]

52. Zhou, Y.; Yang, Z.Y.; Tang, R.C. Bioactive and UV protective silk materials containing baicalin—The multifunctional plant extract from Scutellaria baikalensis Georgi. Mater. Sci. Eng. C Mater. Bioul. Appl. 2016, 67, 336–344. [CrossRef]

53. Cui, L.; Wang, W.; Luo, Y.; Ning, Q.; Xia, Z.; Chen, J.; Feng, L.; Wang, H.; Song, J.; Tan, X.; et al. Polysaccharide from Scutellaria baikalensis Georgi ameliorates colitis via suppressing NF-kappaB signaling and NLRP3 inflammasome activation. Int. J. Biol. Macromol. 2019, 132, 393–405. [CrossRef]

54. Yoon, J.J.; Jeong, J.W.; Choi, E.O.; Kim, M.J.; Hwang-Bo, H.; Kim, H.J.; Hong, S.H.; Park, C.; Lee, D.H.; Choi, Y.H. Protective effects of Scutellaria baikalensis Georgi against hydrogen peroxide-induced DNA damage and apoptosis in HaCaT human skin keratinocytes. EXCLI J. 2017, 16, 426–438. [CrossRef]

55. Jin, B.R.; Chung, K.S.; Kim, H.J.; An, H.J. Chinese Skullcap (Scutellaria baikalensis Georgi) inhibits inflammation and proliferation on benign prostatic hyperplasia in rats. J. Ethnopharmacol. 2019, 235, 481–488. [CrossRef][PubMed]

56. Liu, S.; Xiao, H.; Xiong, Y.; Chen, J.; Wu, Q.; Wen, X.; Jiang, Z.; Wang, L. Effects of Fermented Feed on the Growth Performance, Intestinal Function, and Microbiota of Piglets Weaned at Different Age. Front. Vet. Sci. 2022, 9, 841762. [CrossRef][PubMed]
Animals 2022, 12, 2418

57. Yan, J.; Zhou, B.; Xi, Y.; Huan, H.; Li, M.; Yu, J.; Zhu, H.; Dai, Z.; Ying, S.; Zhou, W.; et al. Fermented feed regulates growth performance and the cecal microbiota community in geese. Poult. Sci. 2019, 98, 4673–4684. [CrossRef]

58. Yeh, H.S.; Weng, B.C.; Lien, T.F. Effects of Chinese traditional herbal medicine complex supplementation on the growth performance, immunity and serum traits of pigs. Anim. Sci. J. 2011, 82, 747–752. [CrossRef]

59. Li, M.; Chen, Z.; Liang, H.; Bai, J.; Zhao, E.; Yi, Z.; Qu, M.; Xu, L. Effects of compound Chinese herbal medicine on growth performance, nutrient apparent digestibility and blood parameters in finishing pigs (in Chinese). China Anim. Husb. Vet. Med. 2019, 46, 1636–1643.

60. Chen, Y.; Sun, Y.; Huang, D.; Wang, D. Effect of Chinese medicine herbal synthiotics on production capability and apparent digestibility of feed on weaned piglets (in Chinese). Acta Ecol. Anim. Domestica 2009, 30, 38–42.

61. Zhou, H.; Xu, R.; Zhang, H.; Su, Y.; Zhu, W. Swine gut microbiota and its interaction with host nutrient metabolism. Nutrients 2019, 11, 2624. [CrossRef] [PubMed]

62. Tett, A.; Pasolli, E.; Masetti, G.; Ercolini, D.; Segata, N. Prevotella diversity, niches and interactions with the human host. Front. Cell. Infect. Microbiol. 2020, 10, 360. [CrossRef]

63. Wu, X.; Guo, C.; Wang, Z.; Peng, Z.; Bai, X.; Li, Y.; Zou, H. Microbiology fermented feed: Effects on performance and nutrient apparent digestibility of lactating dairy cows (in Chinese). Chin. J. Anim. Nutr. 2014, 26, 2296–2302.

64. Wang, J.; Feng, W.; Tang, F.; Ao, H.; Peng, C. Gut microbial transformation, a potential improving factor in the therapeutic activities of four groups of natural compounds isolated from herbal medicines. Fitoterapia 2019, 138, 104293. [CrossRef] [PubMed]

65. An, X.; Bao, Q.; Di, S.; Zhao, Y.; Zhao, S.; Zhang, H.; Lian, F.; Tong, X. The interaction between the gut microbiota and herbal medicines. Biomed. Pharmacother. 2019, 118, 109252. [CrossRef] [PubMed]

66. Zou, J.; Li, W.; Wang, G.; Fang, S.; Cai, J.; Wang, T.; Zhang, H.; Liu, P.; Wu, J.; Ma, Y. Hepatoprotective effects of Huangqi decoction (Astragali Radix and Glycyrrhizae Radix et Rhizoma) on cholestatic liver injury in mice: Involvement of alleviating intestinal microbiota dysbiosis. J. Ethnopharmacol. 2021, 267, 113544. [CrossRef] [PubMed]

67. Zheng, Y.; Gou, X.; Zhang, L.; Gao, H.; Wei, Y.; Yu, X.; Pang, B.; Tian, J.; Tong, X.; Li, M. Interactions between gut microbiota, host, and herbal medicines: A review of new insights into the pathogenesis and treatment of type 2 diabetes. Front. Cell. Infect. Microbiol. 2020, 10, 360. [CrossRef]

68. Long, C.X.; Wu, J.Q.; Tan, Z.J.; Wang, S.P. Different Intestinal Microbiota with Growth Stages of Three-Breed Hybrid Pig. Biomed. Res. Int. 2022, 2022, 5603451. [CrossRef]

69. Qi, K.; Men, X.; Wu, J.; Deng, B.; Xu, Z. Effects of Growth Stage and Rearing Pattern on Pig Gut Microbiota. Curr. Microbiol. 2022, 79, 136. [CrossRef]

70. Zhang, H.Y.; Tian, J.X.; Lian, F.M.; Li, M.; Liu, W.K.; Zhen, Z.; Liao, J.Q.; Tong, X.L. Therapeutic mechanisms of traditional Chinese medicine to improve metabolic diseases via the gut microbiota. Biomed. Pharmacother. 2021, 133, 110857. [CrossRef]

71. Kelly, C.J.; Zheng, L.; Campbell, E.L.; Saeedi, B.; Scholz, C.C.; Bayless, A.J.; Wilson, K.E.; Glover, L.E.; Kominsky, D.J.; Magnuson, A.; et al. Crosstalk between microbiota-derived short-chain fatty acids and intestinal epithelial HIF augments tissue barrier function. Cell Host Microbe 2015, 17, 662–671. [CrossRef]

72. Tett, A.; Pasolli, E.; Masetti, G.; Ercoli, D.; Segata, N. Prevotella diversity, niches and interactions with the human host. Nat. Rev. Microbiol. 2019, 17, 585–599. [CrossRef]

73. Park, S.J.; Kim, J.; Lee, J.S.; Rhee, S.K.; Kim, H. Characterization of the fecal microbiome in different swine groups by high-throughput sequencing. Anaerobe 2014, 28, 157–162. [CrossRef]

74. Rajilic-Stojanovic, M.; Shanahan, F.; Guarner, F.; de Vos, W.M. Phylogenetic analysis of dysbiosis in ulcerative colitis during remission. Inflamm. Bowel Dis. 2013, 19, 481–488. [CrossRef] [PubMed]

75. Jo, H.E.; Kwon, M.S.; Whon, T.W.; Kim, D.W.; Yun, M.; Lee, J.; Shin, M.Y.; Kim, S.H.; Choi, H.J. Alteration of gut microbiota after antibiotic exposure in finishing swine. Front. Microbiol. 2021, 12, 596002. [CrossRef] [PubMed]

76. Wang, H.; Xu, R.; Zhang, H.; Su, Y.; Zhu, W. Swine gut microbiota and its interaction with host nutrient metabolism. Anim. Nutr. 2020, 6, 410–420. [CrossRef]

77. Ye, J.; Zhao, Y.; Chen, X.; Zhou, H.; Yang, Y.; Zhang, X.; Huang, Y.; Zhang, N.; Lui, E.M.K.; Xiao, M. Pu-erh tea ameliorates obesity and modulates gut microbiota in high fat fed mice. Food Res. Int. 2021, 144, 110360. [CrossRef] [PubMed]

78. Hashemi, Z.; Foushe, J.; Im, H.S.; Chan, C.B.; Willing, B.P. Dietary pea fiber supplementation improves glycemia and induces changes in the composition of gut microbiota, serum short chain fatty acid profile and expression of mucins in glucose intolerant rats. Nutrients 2017, 9, 1236. [CrossRef]

79. Liu, Y.; Li, T.; Alam, A.; Ren, D.; Zhao, Y.; Yang, X. Regulatory effects of stachyose on colonic and hepatic inflammation, gut microbiota dysbiosis, and peripheral CD4(+)* T cell distribution abnormality in high-fat diet-fed mice. J. Agric. Food Chem. 2019, 67, 11665–11674. [CrossRef]

80. Niu, Q.; Li, P.; Hao, S.; Zhang, Y.; Kim, S.W.; Li, H.; Ma, X.; Gao, S.; He, L.; Wu, W.; et al. Dynamic distribution of the gut microbiota and the relationship with apparent crude fiber digestibility and growth stages in pigs. Sci. Rep. 2015, 5, 9938. [CrossRef]