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# Dietary coated essential oil and organic acid mixture supplementation improves health of broilers infected with avian pathogenic *Escherichia coli*


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## ABSTRACT

Colibacillosis caused by avian pathogenic *Escherichia coli* (APEC) is a very prevalent disease in poultry farms in China. The exploration of effective non-antibiotic substances is of great significance for the control of APEC infections. This experiment evaluated the efficacy of coated essential oil and organic acid (EOA) supplementation to prevent *E. coli* O78 infection in broiler chickens. A total of 288 one-day-old male broiler chicks were randomly distributed into 4 groups with 6 replicates per group. Chickens were fed a diet either supplemented with EOA (500 mg/kg feed) or not, and either uninfected or infected with *E. coli* O78 intratracheally. Results showed that *E. coli* O78 infection reduced body weight gain, increased mortality and the ratio of feed to gain along with cecal and liver *E. coli* load, damaged gut mucosa, induced local and systemic inflammation, and altered cecal microbial composition, diversity and function ( $P < 0.05$ ). Supplemental EOA improved feed conversion efficiency, lowered gross lesion scores and cecal *E. coli* population, enhanced intestinal goblet cells and serum IgG concentration, and tended to decrease serum IL-12 production ( $P < 0.05$ ). Essential oil and organic acid addition downregulated *IFN- $\gamma$*  mRNA, tended to decrease mucin-2 mRNA levels while upregulating *IL-10* mRNA, and tended to increase *ZO-1* gene expression in the jejunum of infected birds at 7 d after *E. coli* O78 challenge ( $P < 0.05$ ). The 16S rRNA gene sequencing indicated that both EOA addition and *E. coli* O78 challenge altered the diversity and composition of the cecal microbiota community. Furthermore, infected birds fed EOA showed decreased Bacteroidetes and genus *Lactobacillus* abundance compared with the infected control. LEfSe analysis showed that Firmicutes, Ruminococcaceae, Clostridiales, Clostridia, *Lactobacillus*, Lactobacillaceae, and *cc-115* were enriched in the non-infected but EOA-treated group ( $P < 0.05$ ). Collectively, dietary EOA supplementation could mildly alleviate *E. coli*-induced gut injury and inflammation.

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## 1. Introduction

Avian colibacillosis caused by avian pathogenic *Escherichia coli* (APEC) is the most prevalent and detrimental bacterial diseases in the poultry industry (Dho-moulin et al., 1999; Gibbs et al., 2004). In clinical cases of avian colibacillosis, the most frequently detected serotypes of APEC are O1:K1, O2:K1, O5, O8, O15, O35, O53, O55, O150 and O78:K80 (McPeake et al., 2005; Cordoni et al., 2016). APEC infection usually causes colibacillosis, a disease that can be localized or systemic, with various clinical symptoms ranging from

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respiratory tract infection stress, loose white or watery droppings to swollen head syndrome (Dho-moulin et al., 1999). Furthermore, necropsy has revealed that chickens with colibacillosis often exhibit typical lesions including perihepatitis, airsacculitis, pericarditis, egg peritonitis, salpingitis, coligranuloma, omphalitis, cellulitis, embryonic death, septicemia and osteomyelitis/arthritis in poultry (Dziva and Stevens, 2008). As a result, mild and moderate chronic APEC infection often causes reduced feed intake, decreases live weight, egg production and hatching rates, causes poor feed conversion rates, and increases contamination of carcasses at slaughter and thus risks human health (Dho-moulin et al., 1999; Collingwood et al., 2014; Kemmett et al., 2014). Severe and acute APEC infection is usually responsible for higher morbidity and mortality in young chickens and older birds as well as decreased production efficiency (Kemmett et al., 2014; Swelum et al., 2021). Furthermore, APEC can affect all species of poultry in all types of production systems. Colibacillosis in broilers is a critically important disease, which primarily affects broiler chickens between the ages of 3 and 6 wk and is considered the most important cause of growth depression, morbidity and mortality increase, leading to considerable economic losses every year in the global poultry industry (Dho-moulin et al., 1999; Abu-Basha et al., 2012; Zhang et al., 2012; Kemmett et al., 2014). Colibacillosis in broilers often occurs during respiratory stress caused by infection with *Mycoplasma gallisepticum* or viral agents such as infectious bronchitis, Newcastle disease and avian influenza. In addition, immunosuppressive disease (infectious bursal disease) and environmental stresses, such as overcrowding or high levels of dust and ammonia entering through oral and respiratory routes, also lead to APEC systemic infection in broiler chickens (Dho-moulin et al., 1999).

In order to reduce morbidity and mortality related to APEC systemic infection, antimicrobial drugs such as aminoglycosides, amoxicillin, blactams, cephalosporins, florfenicol, fluoroquinolones and tetracyclines, as well as the control of environmental contaminants are often used to manage colibacillosis infection and outbreak. The continuous or excessive use of these antimicrobial drugs in poultry, however, has contributed to the emergence of antimicrobial drug-resistant APEC populations in China and other countries (Zhang et al., 2013; Chen et al., 2014; Zhuang et al., 2014; Ceccarelli et al., 2020; Kim et al., 2020). This phenomenon not only poses new challenges to the rational and effective use of antibiotics, but also brings many difficulties to the prevention and control of APEC infection. Additionally, recent mandated reduction or bans in antimicrobial use in the poultry industry in some countries have impeded the treatment of avian colibacillosis. Consequently, APEC systemic infection incidence and mortality rates have increased gradually. Therefore, the exploration of effective and safe anti-infective substances and vaccines, which carry low risk of drug resistance, is of great significance and is valuable for the control of APEC infections (Swelum et al., 2021).

Essential oils (EO) from plants and Chinese herbal medicines are gathering increased attention and can be utilised as promising antibiotic alternatives due to their effective antimicrobial, antiviral, anti-coccidial, anti-fungal, anti-inflammatory, anti-oxidative and immunomodulative capacities (reviewed by Bakkali et al., 2008; Yang et al., 2015; Zeng et al., 2015; El-Shall et al., 2020). In vitro studies have demonstrated that several plant-derived EO, such as thyme, carvacrol, cinnamaldehyde and citral, could inhibit or kill Gram-negative and Gram-positive bacteria including *Salmonella*, *E. coli*, *Campylobacter* and *Clostridium perfringens*, without harming beneficial bacteria like *Lactobacillus* spp. (Friedman et al., 2002, 2004; Burt, 2004; Mitsch et al., 2004; Jerzsele et al., 2012; Mathlouthi et al., 2012; Calo et al., 2015; Do et al., 2015; Lopez-Romero et al., 2015; Yang et al., 2015). Broiler chicken studies have indicated that several essential oil products with different

kinds of EO components could improve growth performance and gut health (Cross et al., 2007; Khattak et al., 2014; Hashemipour et al., 2015; Pirgozliev et al., 2015; Peng et al., 2016; Adaszyńska-Skwirzyńska and Szczerbińska, 2018; Chowdhury et al., 2018; Liu et al., 2018; Wang et al., 2019), alleviate negative effects caused by pathogenic *Salmonella* (Alali et al., 2013), *E. coli* (Liu et al., 2018), *C. perfringens* (Mitsch et al., 2004; Jerzsele et al., 2012; Du et al., 2015, 2016; Yin et al., 2017) and coccidiosis (Mohiti-Asli and Ghanaatparast-Rashti, 2015; Gordillo Jaramillo et al., 2021).

Organic acids (OA) also could be used as antibiotic alternatives and feed preservatives due to their anti-microbial abilities. Some studies have demonstrated that OA supplementation could improve growth performance, feed conversion rate and feed protein utilization of broiler chickens (Nava et al., 2009; Adil et al., 2011), possibly via suppressing the growth of intestinal acid-intolerant potential pathogens such as *E. coli*, *Salmonella* and *C. perfringens*, increasing intestinal digestive enzyme activity and improving intestinal morphological structure (Gharib Naseri et al., 2012). Furthermore, adding organic acids in drinking water or feed could provide young chickens with protection against *E. coli*, *Salmonella*, *C. perfringens* and *Campylobacter* infection (Chaveerach et al., 2004; Van Immerseel et al., 2006; Dittoe et al., 2018; Lu et al., 2021).

Interestingly, previous studies have indicated that dietary essential oil and organic acid (EOA) supplementation not only showed synergistic beneficial effects on growth performance and gut health, but also exhibited higher efficacy in controlling harmful intestinal bacterial infection such as *E. coli*, *Salmonella* spp. and *C. perfringens* (Giannenas et al., 2014; Aristimunha et al., 2016; Basmacioğlu-Malayoglu et al., 2016; Pham et al., 2020) compared with individual addition. The EOA mixture is a novel blend of coated EOA compounds containing thyme 4%, carvacrol 4%, hexanoic acid 0.5%, benzoic acid 3.5%, butyric acid 0.5% and carrier. However, the protective mechanism of dietary inclusion of the novel EOA mixture in broiler chickens challenged with APEC is unclear. Therefore, the purpose of this study was to evaluate whether adding EOA could protect broiler chickens against *E. coli* O78 infection through determining growth performance, immune response and gut health.

## 2. Materials and methods

### 2.1. Animal ethics statement

All study procedures were approved by the Animal Care and Use Committee of China Agricultural University (permit number: SYXK 2019-0031) and were in accordance with the Beijing Guidelines for Experimental Animals established by the Ministry of Science and Technology (Beijing, China). All efforts were made to minimize the suffering of the animals.

### 2.2. Experimental design, birds and diets

A 2 × 2 completely randomized factorial design was used to evaluate the efficacy of EOA supplementation (0 or 500 mg/kg of diet) and 2 levels of *E. coli* challenge (challenged or unchallenged). A total of 228 one-day-old male (Arbor Acres) broiler chicks were obtained from a company hatchery. After arrival, all chicks were weighed and randomly allocated to 4 groups. There were 6 cages each group with 12 birds per cage per replicate. The treatment groups were as follows: (1) negative control group (**A**, neither EOA treatment nor *E. coli* infection), (2) the EOA-treated group (**B**, EOA treatment at 500 mg/kg of feed but without *E. coli* infection), (3) *E. coli*-infected control group (**K**, *E. coli* infection but without EOA treatment), and (4) the EOA-treated and *E. coli*-infected group (**L**,

both EOA treatment and *E. coli* infection). The EOA product contained vegetable (palm) oil coated thyme 4% and carvacrol 4%, hexanoic acid 0.5%, benzoic acid 3.5%, butyric acid 0.5% and carrier which was provided by Menon Animal Nutrition Technology Co. Ltd, Shanghai, China. To avoid cross-contamination, the uninfected birds and *E. coli*-infected birds were reared in 2 separate areas equipped with three-tiered battery cages with a raised wire-netted floor. The room temperature was maintained at 32 to 34 °C during the first 3 d post-hatch followed by gradual reduction to reach a final room temperature of 22 to 24 °C. The birds received continuous light for the first 3 d (24 h) and then maintained under 23 h/day. In addition, the chickens were vaccinated against Newcastle disease virus (NDV) and infectious bronchitis virus (IBV) on d 7 and 22, respectively; and against infectious bursal disease virus (IBDV) according to the routine immunization program via drinking water on d 12 and 26. All birds were allowed ad libitum access to feed and water throughout the study. Based on the treatments assigned, chickens were fed either antibiotic-free and coccidiostat-free pellet basal diets which were formulated to meet or exceed the National Research Council (1994) requirements for starter (d 1 to 21) and grower (d 22 to 42) periods. The composition of the basal diet and nutrient levels are presented in Table 1.

### 2.3. *Escherichia coli* O78 culture and challenge protocol

The *E. coli* O78 strain (CVCC1418) used in this study was purchased from the Chinese Veterinary Culture Collection Center, which was isolated from the heart of chicken with septicemia signs. The *E. coli* O78 was overnight cultured in Luria–Bertani broth (LB) at 37 °C, and then enumerated by spread-plating the appropriate dilutions in duplicate on tryptose phosphate agar plates. The birds in different treatments were fed the corresponding diet for the first 21 d. On d 22, birds in the K and L groups were intratracheally

challenged with 0.5 mL of *E. coli* O78 suspended in LB at a concentration of  $10^6$  CFU/mL using a polyethylene tube attached to a syringe, as previously described (Kariyawasam et al., 2004; Antão et al., 2008). The birds in the A and B groups were administered the same amount of sterile solution as a control.

### 2.4. Growth performance

Dead birds were recorded daily and mortality rates was calculated at every rearing stage. Each replicate cage had body weight and feed intake (FI) measured on d 1, 21 and 42. Average daily gain (ADG), average daily feed intake (ADFI) and feed conversion ratio (FCR) were calculated and corrected for mortality rate.

### 2.5. Sample collection

At 7 d post-*E. coli* infection, one bird per replicate was randomly selected, weighed, then blood samples were taken from the wing vein and centrifuged ( $3,000 \times g$ , 10 min) at 4 °C, and then the serum was harvested and stored at –20 °C until analysis. The birds were euthanized by cervical dislocation. The middle intestinal sections of the jejunum were cut out (approximately 1 cm), carefully washed with sterile saline, then put into a sterile tube and immediately snap-frozen in liquid nitrogen solution and stored at –80 °C until subsequent analysis for mRNA expression. Another jejunal sample (approximately 2 cm) was rinsed in 0.9% physiological saline and stored in 4% paraformaldehyde buffer solution for later morphological analysis. Liver samples and cecal samples from each bird were aseptically collected into sterile tubes, then immediately snap-frozen in liquid nitrogen solution, stored at –40 °C and moved to the laboratory for microbial culture and microbial 16S rRNA analysis.

### 2.6. Gross lesion examination

At necropsy, tissues were scored for signs of inflammation and lesions in the air sac (0, normal; 1, slight edema; 2, slight diffuse thickening and neovascularization with slight fibrinous exudate; 3, moderate fibrinous exudate; and 4, severe extensive exudate), heart and pericardium (0, normal; 1, vascularization, opacity, cloudy fluid in the pericardial cavity; 2, acute pericarditis), and liver (0, normal; 1, mild fibrinous exudate; 2, severe perihepatitis), as described previously (Peighambari et al., 2000). Scores for heart plus pericardium and liver were combined for final analysis.

### 2.7. Histomorphological structure and goblet cells analysis of the jejunum

Villous height (VH), crypt depth (CD) and goblet cell analysis were performed as previously described (Shao et al., 2013).

### 2.8. Determination of bacterial concentration in the liver and cecal contents

Bacterial concentration in the liver and cecal contents was measured as previously described (Wu et al., 2018). Briefly, 6 birds per group were chosen at random for microbial analyses. The selected broilers were weighed and euthanized, and the liver and cecal contents were aseptically removed. The samples were then diluted 10-fold by weight in 0.90% sterile physiological saline and were mechanically massaged for 1 min. All samples were serially diluted to appropriate levels for plating. An aliquot from each sample (100 µL) was spiral plated on the following: EMB agar for the isolation and enumeration of total coliforms, *Lactobacillus* selection MRS agar for the enumeration of *Lactobacillus* spp.

**Table 1**  
Ingredients and nutrient levels of the basal diet (as-fed basis, %).

Item	1 to 21 d	22 to 42 d
<b>Ingredients</b>		
Corn (7.8% CP)	39.70	57.0
Wheat powder (15.3% CP)	0	5
Wheat (13.8% CP)	19.0	0
Soybean meal (46.0% CP)	33.0	30.0
Soybean oil	4.00	4.40
Limestone-calcium carbonate	1.50	1.50
Calcium hydrogen phosphate	1.50	1.36
Phytase	0.02	0.03
DL-Methionine (98%)	0.27	0.19
L-Lysine HCl (78%)	0.20	0.11
Sodium chloride	0.30	0.30
Vitamin Premix <sup>1</sup>	0.03	0.03
Mineral Premix <sup>2</sup>	0.20	0.20
Choline chloride (50%)	0.25	0.15
Ethoxyquin (33%)	0.05	0.03
Total	100	100
<b>Calculated nutrient levels</b>		
Metabolizable energy, MJ/kg	12.60	12.98
Crude protein	21.37	19.27
Calcium	0.99	0.93
Available phosphorus	0.45	0.43
Lysine	1.20	1.05
Methionine	0.57	0.46
Methionine + Cysteine	0.90	0.78

<sup>1</sup> Vitamin premix provided per kilogram of diet: vitamin A (retinyl acetate), 12,500 IU; vitamin D<sub>3</sub>, 2,500 IU; vitamin E, 30 mg; vitamin K<sub>3</sub>, 3.0 mg; vitamin B<sub>1</sub>, 3.0 mg; vitamin B<sub>2</sub>, 6.0 mg; vitamin B<sub>6</sub>, 6 mg; vitamin B<sub>12</sub>, 0.025 mg; biotin, 0.30 mg; folic acid, 1.25 mg; nicotinic acid, 50 mg; D-pantothenic acid, 12 mg.

<sup>2</sup> Mineral premix provided per kilogram of diet: iron, 80 mg; copper, 8 mg; manganese, 100 mg; zinc, 80 mg; iodine, 0.35 mg; selenium, 0.15 mg.

## 2.9. ELISA for the measurement of immunoglobulins and cytokines

Immune globulin levels such as immunoglobulins (IgG, IgM and IgA), along with cytokines like interleukins (IL-2, IL-4, IL-6), TNF- $\alpha$  and IFN- $\gamma$  were measured in the serum with the use of enzyme-linked immunosorbent assay (ELISA) kits particularly for chickens (R&D Systems, Minneapolis, MN, United States) according to the instructions of the manufacturer.

## 2.10. Quantitative real-time PCR analysis

Trizol reagent (Invitrogen Life Technologies, Carlsbad, CA) was used to isolate total RNA of jejunal mucosa (about 60 mg) according to the manufacturer's instructions. An agarose gel was used to assess the RNA quality. The concentration and purity of the extracted RNA was assessed using a Nanodrop-2000 spectrophotometer (260 and 260/280 nm respectively) (Thermo Fisher Scientific, Waltham, MA, USA). Then, Primer Script RT reagent Kit (TaKaRa Bio Inc.) was used to reverse transcribe RNA to complementary DNA, which was stored at  $-20^{\circ}\text{C}$  until analysis. The target

gene sequences and primers for the reference gene are shown in Table 2. Expressions of toll-like receptor (TLR) signal pathway-related genes, tight junction proteins, growth factors and mucin-2 and housekeeping gene  $\beta$ -actin were quantified with SYBR Premix Ex Taq kits (TaKaRa) on the Applied Biosystems 7500 Fast Real-Time PCR System. The specificity and efficiency of the primer was determined by melt curve analysis. The  $2^{-\Delta\Delta\text{CT}}$  method calculated the levels of mRNA expression of target genes using  $\beta$ -actin as a reference gene (Livak and Schmittgen, 2001).

## 2.11. Metagenomics analysis of gut microbiome

Frozen fecal samples were used for metagenomic studies by analyzing the 16S rDNA, as described previously (Pham et al., 2020). The 16S rRNA gene sequencing was performed using the Illumina MiSeq PE 250 platform (Illumina, Santa Clara, CA) with a MiSeq Reagent Kit at Shanghai Personal Biotechnology Co., Ltd (Shanghai, China). The raw data and QIIME (Quantitative Insights into Microbial Ecology) were filtered and demultiplexed by using the Illumina MiSeq platform (version 1.8.0-dev). Raw tags were obtained by

**Table 2**  
Nucleotide sequences of primers for quantitative real-time PCR assay.

Name	Primer sequence	GenBank accession
Receptors		
TLR-4	F: CCACTATTCCGGTTGGTGGAC R: ACAGCTTCTCAGCAGGCAAT	NM_001030693.1
Adaptor proteins		
MyD88	F: GGATGGTGGTCGTCATTTC R: GAGATTTTCCAGTCCTGTCCA	NM_001030962.1
NF- $\kappa$ B	F: TGGAGAAGGCTATGCAGCTT R: CATCTGGACAGCAGTGAGA	NM_205134.1
Pro-inflammatory cytokines		
TNF- $\alpha$	F: CCCCTACCCTGTCCACAA R: TGAGTACTGCGGAGGGTTCAT	NM204267
IL-1 $\beta$	F: CAGCAGCTCAGCGAAGAG R: CTGTGGTGTGCTCAGAATCCA	NM_204524.1
IL-8	F: GGCTTGCTAGGGGAAATGA R: AGCTGACTCTGACTAGGAAACTGT	AJ009800
IFN- $\gamma$	F: AAAGCCGCACATCAACACA R: GCCATCAGGAAGGTTGTTTTTC	NM_205149.1
Ant-inflammatory cytokines		
IL-10	F: CGCTGTACCCTCTCTCA R: CGTCTCTTGATCTGCTTGATG	NM_001004414.2
Tight junctions		
Claudin-1	F: AAGTGCATGGAGGATGACCA R: GCCACTCTGTGCCATACCA	NM_001013611.2
Occludin	F: TCATCGCCTCCATCGTCTAC R: TCTTACTGCGCGTCTCTGG	NM_205128.1
ZO-1	F: TATGAAGATCGTGCCCTCC R: GAGGTCTGCCATCGTAGCTC	XM_015278981.1
Mucin-2	F: AGCGAGATGTTGGCGATGAT R: AAGTTGCCACACAGACCACA	NM_001318434.1
Growth factors		
TGF- $\beta$ 3	F: TGCCGCCAGATGAGCAT R: TGCACATTCCTGCCACTGA	NM_205454.1
IGF-2	F: TGGCTCTGTGGAAACCTAC R: ACTTGGCATGAGATGGCTTC	NM_001030342.2
EGFR	F: ACCAGCCTGCAGAGAATGTA R: CACCATGTTAAGCGCAATGA	NM_205497
GLP-2	F: AAGCTTCCCAGTCTGAACCA R: ATCCTGAGCTCGTCTGCTGT	NM_205260.4
House-keeping genes		
$\beta$ -actin	F: GAGAAATGTGCGTGACATCA R: CCTGAACCTCTCATTGCCA	NM_205518

TLR = toll-like receptor; MyD88 = myeloid differential protein-88; NF- $\kappa$ B = nuclear factor kappa-light-chain-enhancer of activated B cells; TNF- $\alpha$  = tumor necrosis factor  $\alpha$ ; IL = interleukin; IFN- $\gamma$  = interferon  $\gamma$ ; ZO-1 = zonula occludens-1; EGFR = epidermal growth factor receptor; GLP-2 = glucagon like peptide-2; IGF-2 = insulin-like growth factor-2; TGF- $\beta$ 3 = transforming growth factor beta 3.

merging sequence data using FLAST (Caporaso et al., 2011). The sequences and reference operational taxonomic units (OTU; 97% similarity) were clustered and classified by the UCLUST algorithm and QIIME software. Afterwards, MOTHUR software was used to calculate alpha diversity and beta diversity analysis. The alpha diversity indices, including the observed OTU, ACE, Chao 1, Good's coverage, Shannon, and Simpson index were calculated by MOTHUR, and substantial variations between the microbial compositions in the control and EOA-treated chickens were determined via a nonparametric Mann–Whitney U test ranked using the percentage of representation of individual genera. A Venn diagram was created with R software to visualize the shared and unique OTU among samples based on the Silva taxonomic database (Zaura et al., 2009). Beta diversity analysis was performed to investigate the structural variation of microbial communities across samples using UniFrac distance metrics (Lozupone et al., 2007; Ramette, 2007), Partial least squares discriminant analysis (PLS-DA) (Chen et al., 2011) and nonmetric multidimensional scaling (NMDS) analyses. Linear discriminant analysis effect size (LEfSe) was performed to detect differentially abundant taxa across groups using the default parameters (LDA Score > 2.0,  $P < 0.05$ ) (Segata et al., 2011). Phylogenetic Investigation of Communities by Reconstruction of Unobserved States (PICRUSt 1.0.0) was used to predict metagenome functions associated with bacterial communities based on high-quality 16S rRNA sequencing data (Langille et al., 2013). The functions were deduced using Kyoto Encyclopedia of Genes and Genomes (KEGG) annotations for level 2 pathways.

## 2.12. Data analysis

Data regarding growth performance, gut lesion scores, intestinal bacterial concentration and liver *E. coli*, jejunal morphology and goblet cell numbers, blood parameters, the relative expression of mRNA, alpha diversity indices of Shannon and abundance-based coverage estimator (ACE) among the 4 groups were analyzed using one-way ANOVA in SPSS 20.0 (SPSS Inc., Chicago, IL, USA) in a 2 × 2 factorial design. Mean separations were carried out using Duncan's multiple comparison test when interactive effects differed significantly. The Kruskal–Wallis test as well as Benjamini–Hochberg  $P$ -value correction was used to compare the results of phylum and genus abundances.  $P \leq 0.05$  was regarded as significant, while  $0.05 \leq P \leq 0.10$  was regarded as a tendency.

**Table 3**

Effects of dietary coated essential oil and organic mixture supplementation on growth performance of broiler chickens infected with *Escherichia coli* O78.

Item	Experimental design				SEM	Main effect				$P$ -value <sup>1</sup>		
	A	K	B	L		0	500 mg/kg	Non-challenge	Challenged	EOA	Challenged	Interaction effects
Day 1 to 21												
ADG, g/d	41.4	42.4	40.8	42.7	0.36	41.9	41.8	41.1	42.6	0.811	0.074	0.575
ADFI, g/d	59.5	60.0	59.3	60.7	0.43	59.8	60.0	59.4	60.4	0.792	0.337	0.672
FCR, g/g	1.44	1.41	1.45	1.42	0.01	1.43	1.44	1.44 <sup>a</sup>	1.42 <sup>b</sup>	0.294	0.046	0.677
Day 22 to 42												
ADG, g/d	90.7	80.5	89.7	80.5	1.79	85.6	85.1	90.2	80.5	0.927	0.035	0.235
ADFI, g/d	147.4 <sup>a</sup>	144.8 <sup>ab</sup>	127.8 <sup>b</sup>	136.8 <sup>ab</sup>	3.10	146.1	132.3	137.6	140.8	0.027	0.582	0.036
FCR, g/g	1.66 <sup>ab</sup>	1.81 <sup>a</sup>	1.44 <sup>c</sup>	1.71 <sup>ab</sup>	0.06	1.73	1.58	1.55	1.76	0.193	0.098	<0.001
Day 1 to 42												
ADG, g/d	66.1	61.5	65.2	61.6	1.17	63.8	63.4	65.6	61.5	0.896	0.129	0.263
ADFI, g/d	103.4 <sup>a</sup>	102.4 <sup>ab</sup>	93.6 <sup>b</sup>	98.8 <sup>ab</sup>	1.56	102.9	96.2	98.5	100.6	0.031	0.476	0.034
FCR, g/g	1.58 <sup>b</sup>	1.67 <sup>a</sup>	1.44 <sup>c</sup>	1.61 <sup>a</sup>	0.04	1.63	1.52	1.51	1.64	0.216	0.129	<0.001

A = neither infection nor coated essential oil and organic acid mixture addition negative control; K = *E. coli* infection positive control; B = coated essential oil and organic acid mixture addition at 500 mg/kg but without infection; L = both essential oil and organic acid mixture addition at 500 mg/kg and *E. coli* O78 infection; SEM = standard error of the mean; ADG = average daily weight gain; ADFI = average daily feed intake; FCR = feed-to-gain ratio.

<sup>a, b, c</sup> Means within the same row without a common superscript differ significantly ( $P < 0.05$ ).

<sup>1</sup>  $P$ -values represent the main effect of the diet, the main effect of *E. coli* O78 challenge, and the interaction between the dietary treatments and *E. coli* O78 challenge.

## 3. Results

### 3.1. Growth performance

As illustrated in Table 3, dietary EOA addition (B and L) had no significant effects on ADG, ADFI and FCR of broiler chickens except for showing a numerical decrease in mortality rate compared with the un-treated birds (A and B) from 1 to 21 d. However, *E. coli* challenge retarded the growth of chickens with an obvious reduction in ADG ( $P < 0.05$ ) from 22 to 42 d, and remarkably increased mortality rate (Fig. 1A) and tended to increase FCR from 22 to 42 d and 1 to 42 d ( $P < 0.05$ ) relative the non-challenged groups. While birds fed with EOA showed decreased ADFI and numerically reduced mortality rate (Fig. 1A) from 22 to 42 d and 1 to 42 d, compared with the untreated groups ( $P < 0.05$ ). There was a notable interaction effect for ADFI and FCR between EOA supplementation and *E. coli* challenge during the later and entire period ( $P > 0.05$ ). Birds on L group exhibited improved FCR compared with infected control group.

### 3.2. Bacterial concentration of intestinal contents and liver

As shown in Table 4, a significant interaction effect between *E. coli* O78 infection and EOA addition on *E. coli* concentration in cecal chyme ( $P < 0.05$ ). Both broilers on L and B groups displayed lower levels of *E. coli* in cecal chyme ( $P < 0.05$ ) compared with the *E. coli*-challenged birds. In addition, *E. coli* challenge tended to increase *E. coli* concentration in liver ( $P = 0.057$ ).

### 3.3. Gross lesion examination

Intratracheal inoculation of *E. coli* O78 not only caused higher mortality, but also lead to severe airsacculitis, pericarditis and perihepatitis. Fig. 1B illustrates that the highest gross lesion scores were observed in the *E. coli* O78-challenged birds ( $P < 0.05$ ). However, EOA addition notably lowered gross lesion scores of birds on L group compared with the challenged control group ( $P < 0.05$ ). No obvious pathological changes were observed in A and B groups.

### 3.4. Intestinal morphology

Based on the main effect of EOA supplementation, birds fed diets with EOA significantly increased villous height (VH), the villous



**Table 5**Effect of dietary coated essential oil and organic mixture supplementation on jejunal morphology and goblet cell numbers of broiler chickens challenged with *Escherichia coli* O78.

Item	<i>E. coli</i> <sup>1</sup>	Villous height, $\mu\text{m}$	Crypt depth, $\mu\text{m}$	VH:CD ratio	GC cells
EOA, mg/kg					
0	–	439.66	61.93	7.07	23.57 <sup>a</sup>
0	+	377.68	79.94	4.92	19.30 <sup>b</sup>
500	–	507.07	59.00	8.66	23.03 <sup>ab</sup>
500	+	491.23	67.72	7.62	26.30 <sup>a</sup>
SEM		20.36	4.19	0.35	0.81
Main effect					
0		408.67 <sup>b</sup>	70.94	5.99 <sup>b</sup>	21.43
500		499.15 <sup>a</sup>	63.36	8.14 <sup>a</sup>	24.67
Non-challenge		473.37	60.47	7.86 <sup>a</sup>	23.30
Challenge		434.46	73.83	6.27 <sup>b</sup>	22.80
<i>P</i> -value <sup>2</sup>					
EOA		0.025	0.368	<0.001	0.024
Challenged		0.308	0.120	0.001	0.711
EOA $\times$ Challenged		0.542	0.578	0.181	0.010

EOA = coated essential oil and organic mixture; VH:CD ratio = the villus height to crypt depth ratio; GC cells = goblet cell numbers per mm<sup>2</sup>; SEM = standard error of the mean.<sup>a, b</sup> Means within the same column with different superscripts differ significantly ( $P < 0.05$ ).<sup>1</sup> “–”, without *E. coli* O78 challenge; “+”, with *E. coli* O78 challenge.<sup>2</sup> *P*-values represent the main effect of the diet, the main effect of *E. coli* O78 challenge, and the interaction between the dietary treatments and *E. coli* O78 challenge.**Table 6**Effect of dietary coated essential oil and organic mixture supplementation on serum cytokine concentration and immunoglobulin levels of broiler chickens challenged with *Escherichia coli* O78.

Item	<i>E. coli</i> <sup>1</sup>	IL-1 $\beta$ , pg/mL	IL-4, pg/mL	IL-6, pg/mL	IL-8, pg/mL	IL-10, pg/mL	IL-12, pg/mL	IFN- $\gamma$ , pg/mL	TNF- $\alpha$ , pg/mL	IgG, g/L
EOA, mg/kg										
0	–	12.13	25.74	153.57	26.68	46.12	21.42 <sup>b</sup>	29.55	4.06	3.81 <sup>c</sup>
0	+	19.03	53.23	794.79	80.48	47.77	166.52 <sup>a</sup>	454.74	36.41	3.75 <sup>c</sup>
500	–	18.25	17.30	172.17	31.61	59.85	46.86 <sup>b</sup>	73.00	9.66	4.29 <sup>b</sup>
500	+	10.15	36.28	369.97	41.20	56.90	82.81 <sup>ab</sup>	237.61	22.73	5.12 <sup>a</sup>
SEM		2.79	5.12	86.36	7.58	9.12	16.50	51.99	4.80	0.13
Main effect										
0		15.58	39.48	474.18	53.58	46.95	93.97	242.14	20.23	3.78
500		14.20	26.79	271.07	36.41	58.38	64.83	155.30	16.20	4.71
Non-challenge		15.19	21.52 <sup>b</sup>	162.87 <sup>b</sup>	29.15 <sup>b</sup>	52.99	34.14	51.27 <sup>b</sup>	6.86 <sup>b</sup>	4.05
Challenge		14.59	44.75 <sup>a</sup>	582.38 <sup>a</sup>	60.84 <sup>a</sup>	52.33	124.66	346.17 <sup>a</sup>	29.57 <sup>a</sup>	4.44
<i>P</i> -value <sup>2</sup>										
EOA		0.813	0.182	0.176	0.209	0.562	0.267	0.307	0.645	<0.001
Challenged		0.918	0.020	0.009	0.026	0.973	0.002	0.002	0.016	0.018
EOA $\times$ Challenged		0.206	0.649	0.142	0.110	0.907	0.045	0.131	0.278	0.007

TNF- $\alpha$  = tumor necrosis factor  $\alpha$ ; IL = interleukin; IFN- $\gamma$  = interferon  $\gamma$ ; IgG = immunoglobulin G; EOA = coated essential oil and organic mixture; SEM = standard error of the mean.<sup>a, b, c</sup> Means within the same column with different superscripts differ significantly ( $P < 0.05$ ).<sup>1</sup> “–”, without *E. coli* O78 challenge; “+”, with *E. coli* O78 challenge.<sup>2</sup> *P*-values represent the main effect of the diet, the main effect of *E. coli* O78 challenge, and the interaction between the dietary treatments and *E. coli* O78 challenge.

infected control alone at 7 d post-challenge. Furthermore, the highest expression levels of mucin-2 were found in the uninfected and EOA treated group relative to that of the other three groups.

### 3.7. Diversity and composition of cecal microbiota

In this study, we also determined cecal microbial communities of broiler chickens treated with EOA supplementation and challenged with *E. coli* O78. A total of 4,766 OTU were obtained from the cecal contents of all groups through high-throughput sequence analysis of the bacterial 16S rRNA V3–V4 region. Venn diagram analysis illustrated that 4,253 common OTU were shared by all 4 groups, whereas unique OTU varied from 40 to 75 in one group, respectively (Fig. 2).

Results of alpha diversity (Fig. 3) showed that *E. coli* O78 infection remarkably increased Simpson index ( $P = 0.038$ ), and tended to increase Shannon index ( $P = 0.053$ ), while birds fed with EOA significantly increased Chao 1 and ACE indices compared with

nonsupplemented groups ( $P < 0.05$ ). There was a considerable interaction effect on Chao 1 and ACE indices between EOA supplementation and APEC challenge. Challenged birds given EOA had higher Chao 1 and ACE indices compared with the single challenged control. Results of alpha diversity indicated that EOA administration significantly increased richness of the intestinal microbial community, while the intestinal flora was more evenly distributed following *E. coli* O78 infection. Beta diversity of the cecal microbial community indicated that there was no obvious convergence among the 4 clusters of the microbial communities from the 4 groups. In addition, the clusters of the cecal microbial communities from the infected groups (K and L) were separated from those of the non-infected groups (A and B). Furthermore, the non-infected groups clustered closely together while the infected groups were highly dispersed.

Cecal microbial composition analysis indicated that Firmicutes, Bacteroidetes and Proteobacteria dominated the cecal microbial community of chickens (>90%) (Fig. 4). Birds fed with EOA

**Table 7**Effects of dietary coated essential oil and organic mixture supplementation on gene expressions of TLR signaling pathway-related genes in the jejunum of broiler chickens challenged with *E. coli* O78.

Item	<i>E. coli</i> <sup>1</sup>	<i>TLR-4</i>	<i>MyD88</i>	<i>NF-κB</i>	<i>IFN-γ</i>	<i>TNF-α</i>	<i>IL-1β</i>	<i>IL-8</i>	<i>IL-10</i>
EOA, mg/kg									
0	–	1.07	1.02	1.06	1.01	1.00	1.05	1.04	1.07 <sup>b</sup>
0	+	1.92	1.18	1.04	2.74	1.04	1.12	1.38	0.72 <sup>b</sup>
500	–	1.18	1.11	1.16	1.13	1.32	1.02	0.75	0.81 <sup>b</sup>
500	+	1.31	1.22	1.03	1.06	1.13	1.09	1.96	1.84 <sup>a</sup>
SEM		0.10	0.08	0.09	0.09	0.06	0.14	0.32	0.11
Main effect									
0		1.50	1.10	1.05	1.87	1.02	1.09	1.21	0.90
500		1.20	1.17	1.10	1.09	1.23	1.05	1.36	1.33
Non-challenged		1.13	1.07	1.11	1.07	1.16	1.04	0.90	0.94
Challenged		1.66	1.20	1.03	1.90	1.09	1.10	1.67	1.28
<i>P</i> -value <sup>2</sup>									
EOA		0.377	0.663	0.805	0.408	0.095	0.905	0.821	0.055
Challenged		0.032	0.407	0.676	0.016	0.524	0.803	0.013	0.126
Challenged × EOA		0.567	0.877	0.747	<0.01	0.353	0.998	0.237	<0.01

*TLR* = toll-like receptor; *MyD88* = myeloid differential protein-88; *NF-κB* = nuclear factor kappa-light-chain-enhancer of activated B cells; *TNF-α* = tumor necrosis factor α; *IL* = interleukin; *IFN-γ* = interferon γ; SEM = standard error of the mean.

<sup>a, b</sup> Means within the same column with different superscripts differ significantly ( $P < 0.05$ ).

<sup>1</sup> “–”, without *E. coli* O78 challenge; “+”, with *E. coli* O78 challenge.

<sup>2</sup> *P*-values represent the main effect of the diet, the main effect of *E. coli* O78 challenge, and the interaction between the dietary treatments and *E. coli* O78 challenge.

**Table 8**Effects of dietary coated essential oil and organic mixture supplementation on gene expressions of tight junction proteins, growth factors and mucin-2 in the jejunum of broiler chickens challenged with *Escherichia coli* O78.

Item	<i>E. coli</i> <sup>1</sup>	Claudin1	Occludin	<i>ZO-1</i>	<i>TGF-β3</i>	<i>IGF-2</i>	<i>EGFR</i>	<i>GLP-2</i>	Mucin-2
EOA, mg/kg									
0	–	0.82	1.02	1.07 <sup>a</sup>	1.03	1.04	1.02	1.04	1.05 <sup>b</sup>
0	+	0.71	0.52	0.27 <sup>c</sup>	1.81	3.22	1.18	0.95	1.13 <sup>ab</sup>
500	–	0.50	1.10	0.64 <sup>b</sup>	1.18	1.21	1.45	1.26	1.44 <sup>a</sup>
500	+	0.80	0.72	0.48 <sup>bc</sup>	1.51	3.28	1.13	1.25	0.93 <sup>b</sup>
SEM		0.06	0.05	0.05	0.12	0.20	0.11	0.08	0.06
Main effect									
0		0.76	0.77	0.67	1.42	2.13	1.10	1.00	1.09
500		0.65	0.91	0.56	1.34	2.25	1.29	1.25	1.19
Non-challenged		0.66	1.06 <sup>a</sup>	0.86	1.10 <sup>b</sup>	1.13 <sup>b</sup>	1.24	1.15	1.25
Challenged		0.75	0.62 <sup>b</sup>	0.38	1.66 <sup>a</sup>	3.25 <sup>a</sup>	1.16	1.10	1.03
<i>P</i> -value <sup>2</sup>									
EOA		0.415	0.175	0.255	0.745	0.777	0.406	0.112	0.455
Challenged		0.470	<0.001	<0.001	0.035	<0.001	0.728	0.735	0.101
Challenged × EOA		0.125	0.589	0.004	0.373	0.891	0.304	0.789	0.028

*ZO-1* = zonula occludens-1; *EGFR* = epidermal growth factor receptor; *GLP-2* = glucagon like peptide-2; *IGF-2* = insulin-like growth factor-2; *TGF-β3* = transforming growth factor beta 3; EOA = coated essential oil and organic mixture; SEM = standard error of the mean.

<sup>a, b, c</sup> Means within the same column with different superscripts differ significantly ( $P < 0.05$ ).

<sup>1</sup> “–”, without *E. coli* O78 challenge; “+”, with *E. coli* O78 challenge.

<sup>2</sup> *P*-values represent the main effect of the diet, the main effect of *E. coli* O78 challenge, and the interaction between the dietary treatments and *E. coli* O78 challenge.

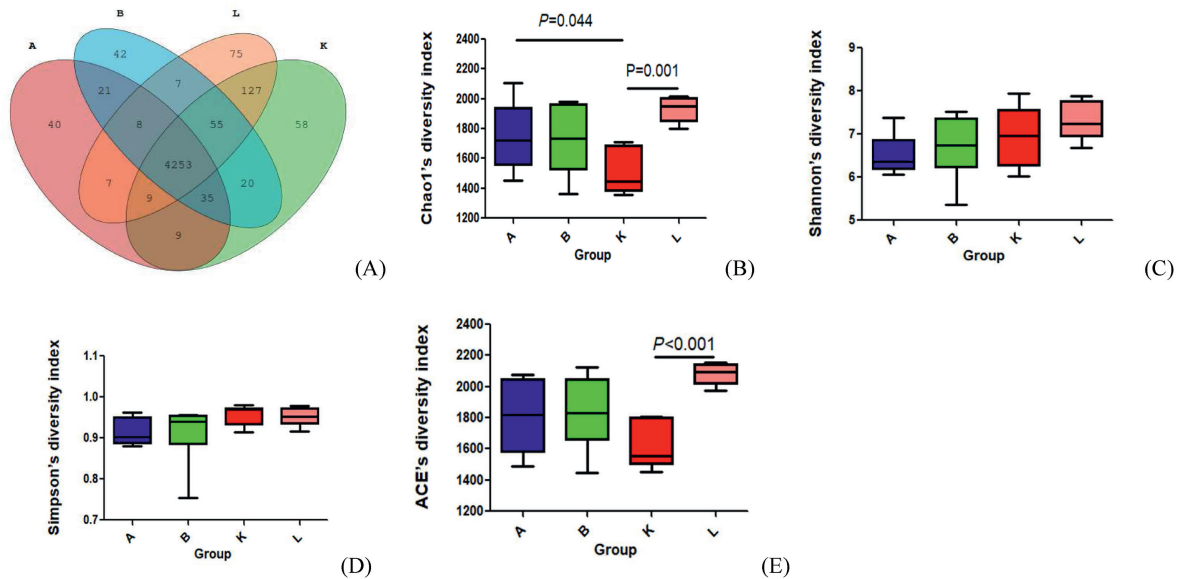
significantly increased relative abundance of Firmicutes compared with the untreated groups, regardless of *E. coli* challenge. Infected birds that received EOA also notably suppressed the enrichment of Bacteroidetes compared with the single infected control ( $P = 0.015$ ). However, *E. coli* infection alone decreased the relative population of Firmicutes ( $P = 0.007$ ) relative to the negative control without EOA addition nor challenge (Fig. 4). Fig. 5 shows relative abundance of the cecal microbial composition at the genus level (Top 10). Results revealed that *Faecalibacterium*, *Ruminococcus*, *Lactobacillus*, *Enterococcus*, *Oscillospira* and *Bilophila* were the predominant genera, followed by *Dorea* and *Coprococcus*. Although there was no significant difference between the negative control group (A) and EOA group (B), the infected birds fed EOA showed a decrease in relative distribution of the genus *Lactobacillus* ( $P = 0.03$ ) compared with the infected birds (K), whereas birds on K group remarkably enriched the relative fraction of the genus *Lactobacillus* and *Oscillospira* (Fig. 5B).

LEfSe analysis was used to determine the statistically different biomarkers between groups. As presented in Fig. 6, significant differences were only observed between groups B and K.

When compared with the B group, *E. coli* challenge alone (K) enriched the relative abundance of *Rikenellaceae*, *Bacteroidales*, *Bacteroidia*, *Bacteroidetes*, *YS2*, *4C0d\_2*, and *Cyanobacteria*, while *Firmicutes*, *Ruminococcaceae*, *Clostridiales*, *Clostridia*, *Lactobacillus*, *Lactobacillaceae*, and *cc-115* were more abundant in the EOA-treated group (B) in contrast to the K group.

### 3.8. Predicted function of cecal microbiota

PICRUSt analysis showed significant distinctive differences between group A and K at KEGG level 2 (Fig. 7). We found that pentose and glucuronate interconversion ( $P = 0.022$ ), and photosynthesis proteins ( $P = 0.057$ ) were suppressed, while folate metabolism, methane metabolism, glycine, serine and threonine metabolism,



**Fig. 2.** The overall description of gut microorganism among the 4 groups. (A) Venn diagram, (B) Chao index, (C) Shannon index, (D) Simpson index, (E) abundance-based coverage estimator (ACE). Treatment groups: A = neither infection nor essential oil and organic acid mixture (EOA) addition control, K = *Escherichia coli* 078 infection control, B = EOA addition but without infection, L = both EOA addition and *E. coli* 078 infection.

MAPK signaling pathway, oxidative phosphorylation and beta-alanine metabolism were overrepresented ( $P = 0.057$ ) in group K compared with the negative control A. However, there was no obvious difference between group K and L through PICRUST analysis ( $P > 0.05$ ).

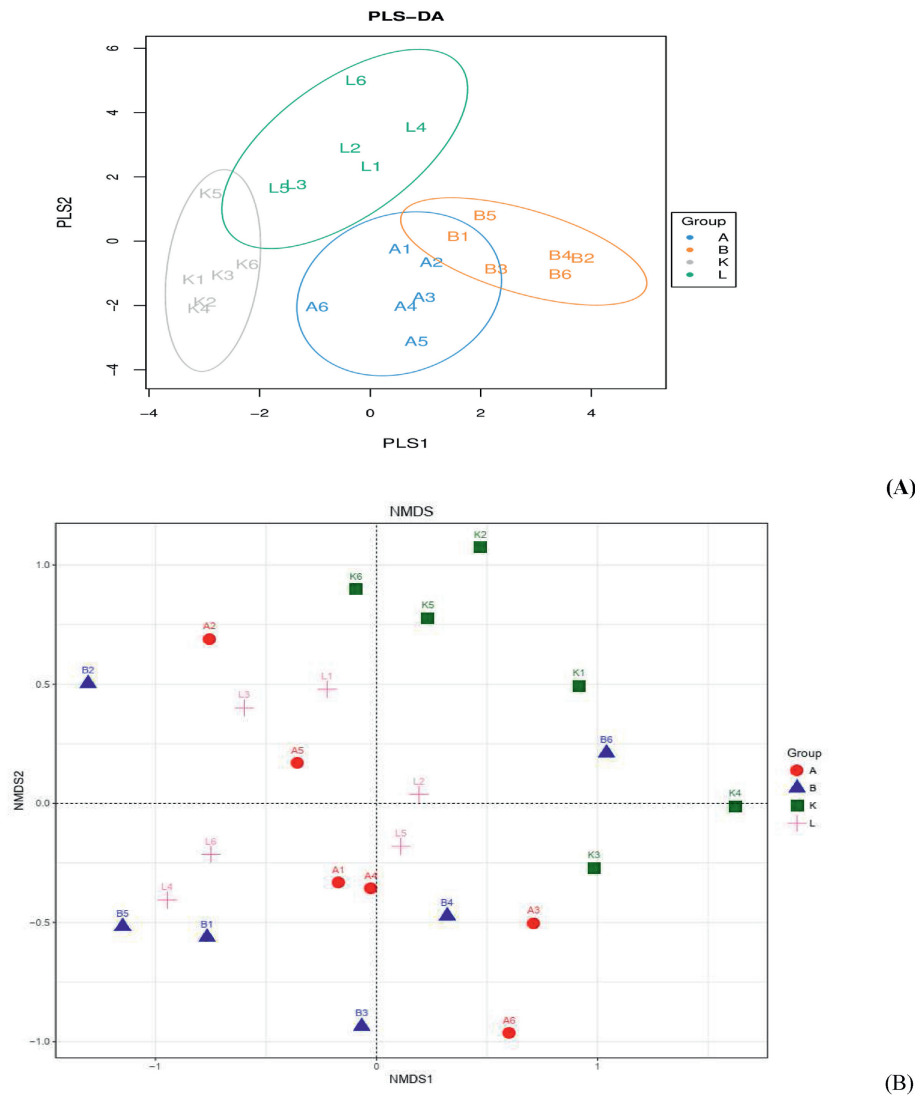
#### 4. Discussion

This study evaluated the preventive efficacy of supplemental EOA on broiler chickens challenged intratracheally with *E. coli* 078. Results from the current study indicated that mortality, feed gain, and gross lesion scores were remarkably increased by intratracheal infection of *E. coli* 078, which was consistent with results of previous studies (Kariyawasam et al., 2004; Horn et al., 2012), suggesting that intratracheal infection of pathogenic *E. coli* 078 can spread from the respiratory tract to various organs, resulting in severe systemic inflammation and negative effects on growth performance, livability and health. However, these negative impacts caused by *E. coli* 078 infection were attenuated by EOA supplementation. Additionally, although challenged birds given EOA showed a numerical reduction in mortality and had no improvement in body weight gain relative to the infected birds alone, EOA administration notably improved feed conversion efficiency, which was attributed to lower feed intake of challenged broiler chickens. In line with our findings, previous studies suggested that dietary essential oils and organic acid blend inclusion resulted in adverse or no effects on performance in chickens (Fascina et al., 2012, 2017), and early-weaned pigs (Manzanilla et al., 2004). Whereas other researches have reported that dietary EOA addition (containing thyme, carvacrol and butyric acid) could improve growth performance, as evidenced by an increase in average body weight gain, or improved FCR, when broiler chickens were either challenged or unchallenged with pathogens (Timbermont et al., 2010; Jerzsele et al., 2012; Abdelli et al., 2020; Stamilla et al., 2020; Stefanello et al., 2020; Pham et al., 2021). Variable results in growth performance of broiler chickens induced by EOA administration were possibly attributed to differences in chemical composition and dosage of EOA used; characteristics of infection pathogen and challenge route; birds' health status; and

the hygiene conditions of the experimental environments. However, our findings suggested that the novel EOA used in the study might elicit mild health-improving effects on broiler chickens intratracheally challenged with *E. coli* 078. Furthermore, this study indicated that dietary EOA supplementation improved feed efficiency of broilers and reduced the mortality after *E. coli* 078 challenge.

Gut morphology, goblet cell numbers, intestinal pathogen load and bacterial translocation are important indirect markers for assessing intestinal health, recovery and functionality (Kuttappan et al., 2015; Celi et al., 2017). The present study found that *E. coli* 078 intratracheal infection could damage gut health, as evidenced by decreased villus height and VH:CD ratio, lower goblet cell numbers, and higher *E. coli* carrier in the cecum and liver, which was similar to our previous observations (Huang et al., 2019). However, feeding EOA lowered *E. coli* counts in cecal chyme, prevented bacterial translocation and improved intestinal morphological structure regardless of *E. coli* 078 infection. Moreover, elevated goblet cell density was also observed in the infected birds treated with EOA relative to the single infected control. Similarly, previous studies from poultry and pig trials also indicated that supplemental EOA reduced the concentration of potential intestinal pathogens including *E. coli*, *Salmonella* and *C. perfringens* (Timbermont et al., 2010; Jerzsele et al., 2012; Cerisuelo et al., 2014; Basmacioğlu-Malayoğlu et al., 2016; Yang et al., 2019; Zhang et al., 2019; Abdelli et al., 2020; Stamilla et al., 2020; Stefanello et al., 2020), and attenuated gut impairment caused by pathogens (Timbermont et al., 2010; Jerzsele et al., 2012; Abdelli et al., 2020; Stamilla et al., 2020; Stefanello et al., 2020). Therefore, our results indicated that feeding EOA could mitigate gut impairment caused by *E. coli* 078 infection. Gut barrier-protecting effects of EOA in this study were possibly associated with the anti-bacterial capacity of EO or OA.

Serum immune parameters, such as cytokines, antibodies and complement proteins, were always used to evaluate systemic immune function or health status of the host (Lee et al., 2013). In this study, *E. coli* 078 infection induced a systemic inflammation response, indicated by higher levels of serum IgG and pro-inflammatory cytokines, such as IL-4, IL-6, IL-8, IL-12, IFN- $\gamma$  and

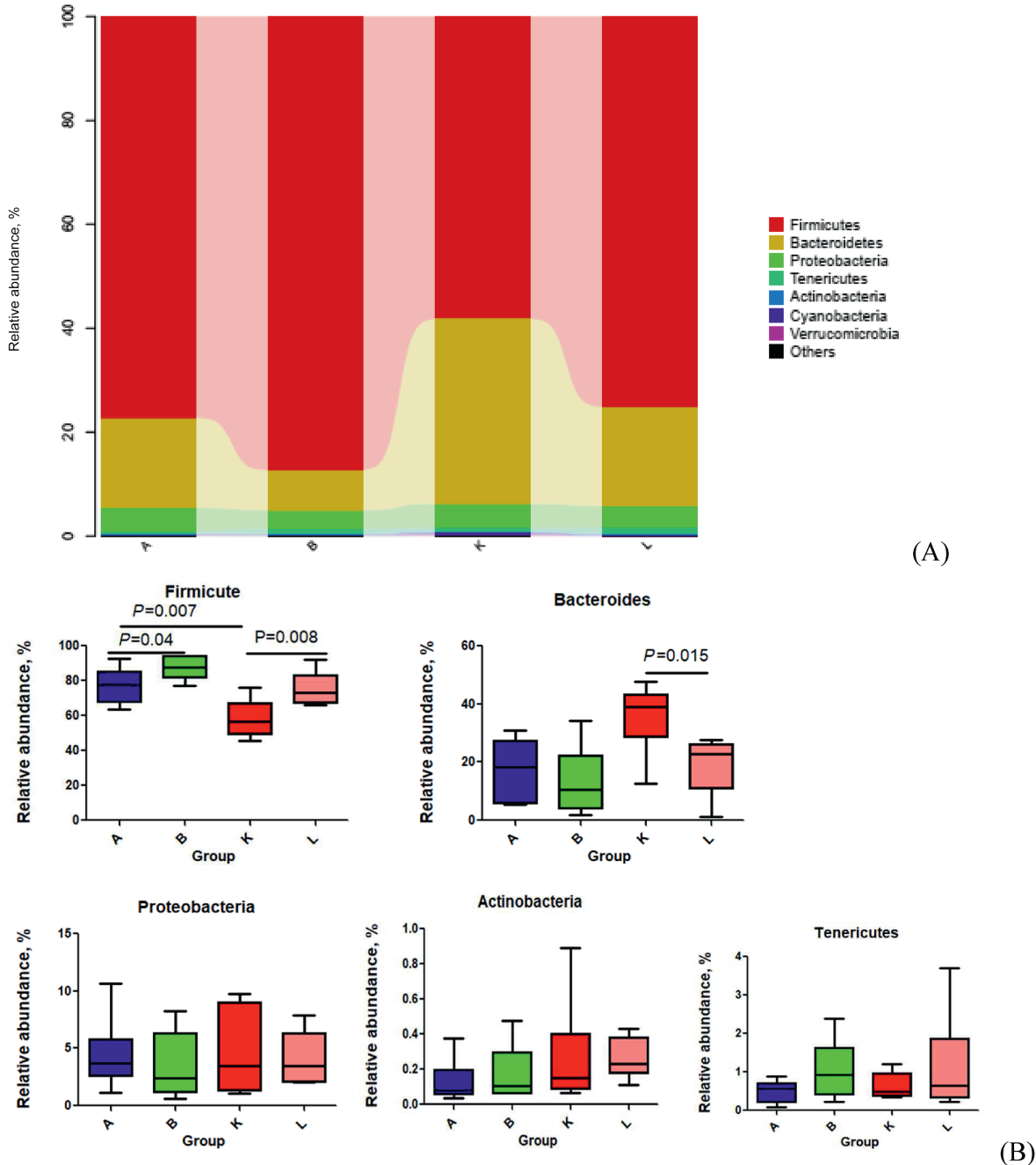


**Fig. 3.** Beta diversity analysis of cecal microbial community. (A) Partial least squares discriminant analysis (PLS-DA). (B) Nonmetric multidimensional scaling (NMDS). Treatment groups: A = neither infection nor essential oil and organic acid mixture (EOA) addition control, K = *Escherichia coli* O78 infection control, B = EOA addition but without infection, L = both EOA addition and *E. coli* O78 infection.

TNF- $\alpha$ . Overproduction of pro-inflammatory cytokines caused by pathogens may lead to damage in the body and gut (McKay and Baird, 1999). Higher gross lesion scores and damaged gut morphological structure caused by *E. coli* O78 infection were observed in this study, which supported our findings. Nevertheless, supplemental EOA significantly increased serum IgG levels and tended to decrease serum IL-12 production compared with the infected groups alone. IgG, secreted by B cells, directly contributes to an immune response by neutralizing of pathogens, toxins and viruses (Song et al., 2018). IL-12 is also involved in pro-inflammatory responses (Vignali and Kuchroo, 2014). Thus, lower pro-inflammatory cytokine IL-12 concentration and higher levels of serum IgG, accompanied by lower bacterial translocation found in the infected birds receiving EOA suggested that EOA supplementation could alleviate systemic inflammatory responses caused by *E. coli* O78 infection.

Tight junctions between intestinal epithelial cells form a barrier with selective permeability that not only allows nutrients, ions and solutes to be transferred but also helps to maintain the integrity of the intestinal mucosa barrier, immune hemostasis and intestinal

health by preventing intestinal microorganisms, antigens and toxins from translocating into tissues (Schneeberger and Lynch, 2004; Turner, 2009; Gil-cardoso et al., 2016; Awad et al., 2017; Lee et al., 2018). Cytokines related to TLR-mediated signaling pathways and the energy status of enterocytes might all have an impact on tight junctions and intestinal epithelial barrier integrity (Capaldo and Nusrat, 2009; Lee, 2015). Pro-inflammatory cytokines have been shown to cause tight junction disruption, which results in increased intestinal permeability, whereas anti-inflammatory cytokines have been found to maintain intestinal integrity (Al-Sadi et al., 2009). In this study, we further examined the alterations in the gut mucosa TLR-mediated immune responses and tight junction proteins in *E. coli*-infected broiler chickens. Results indicated that *E. coli* O78 intratracheal infection induced intestinal inflammation and damaged intestinal barrier function, as evidenced by notable upregulation in gene expression of TLR4 and pro-inflammatory cytokines IFN- $\gamma$  and chemokine IL-8, as well as downregulation in tight junction proteins Occludin and ZO-1 mRNA level in the jejunum. However, feeding EOA remarkably lowered jejunal pro-inflammatory cytokine IFN- $\gamma$  abundance, but

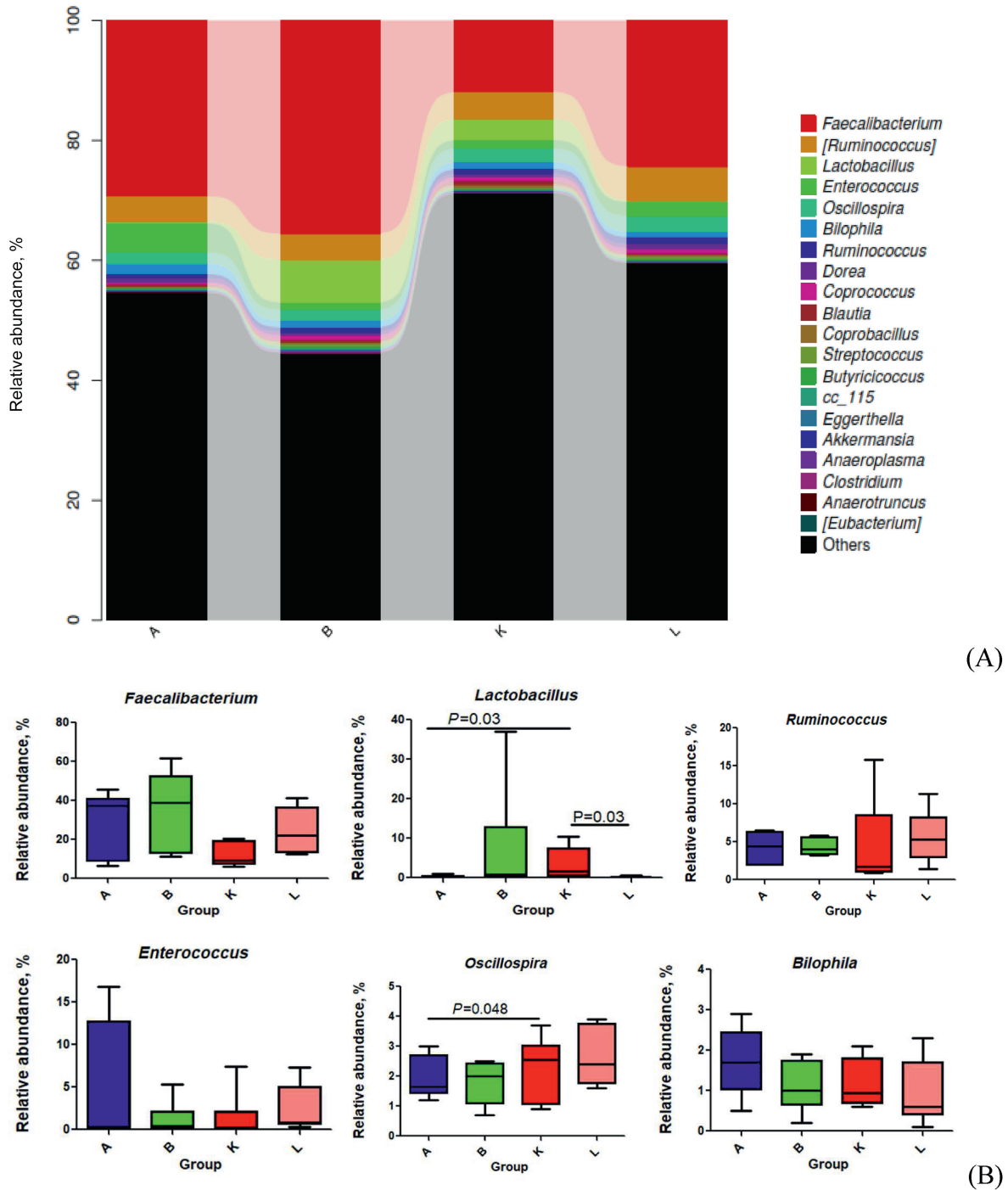


**Fig. 4.** Composition of caecal microbiota of the broiler chickens among the 4 groups at phylum level. (A) Relative abundance of top 8 bacterial phyla. (B) Comparison of the relative abundances of the top 5 bacterial phyla among the 4 groups. Treatment groups: A = neither infection nor essential oil and organic acid mixture (EOA) addition control, K = *Escherichia coli* O78 infection control, B = EOA addition but without infection, L = both EOA addition and *E. coli* O78 infection.

increased anti-inflammatory cytokine IL-10 mRNA levels and tended to increase ZO-1 gene expression of the *E. coli*-infected birds. In agreement with our findings, previous studies have demonstrated that a mixture of supplemental coated essential oils and organic acid (including thyme, carvacrol and butyric acid) could mitigate gut injury caused by oral *C. perfringens* infection (Timbermont et al., 2010; Jerzsele et al., 2012; Abdelli et al., 2020; Pham et al., 2020; Stamilla et al., 2020; Stefanello et al., 2020) through inhibiting intestinal inflammatory responses and improving the energy status of enterocytes. Thus, based on our results, we have suggested that dietary EOA supplementation protected against APEC-induced

intestinal barrier junction injury in broilers, possibly through alleviating intestinal inflammatory responses caused by APEC.

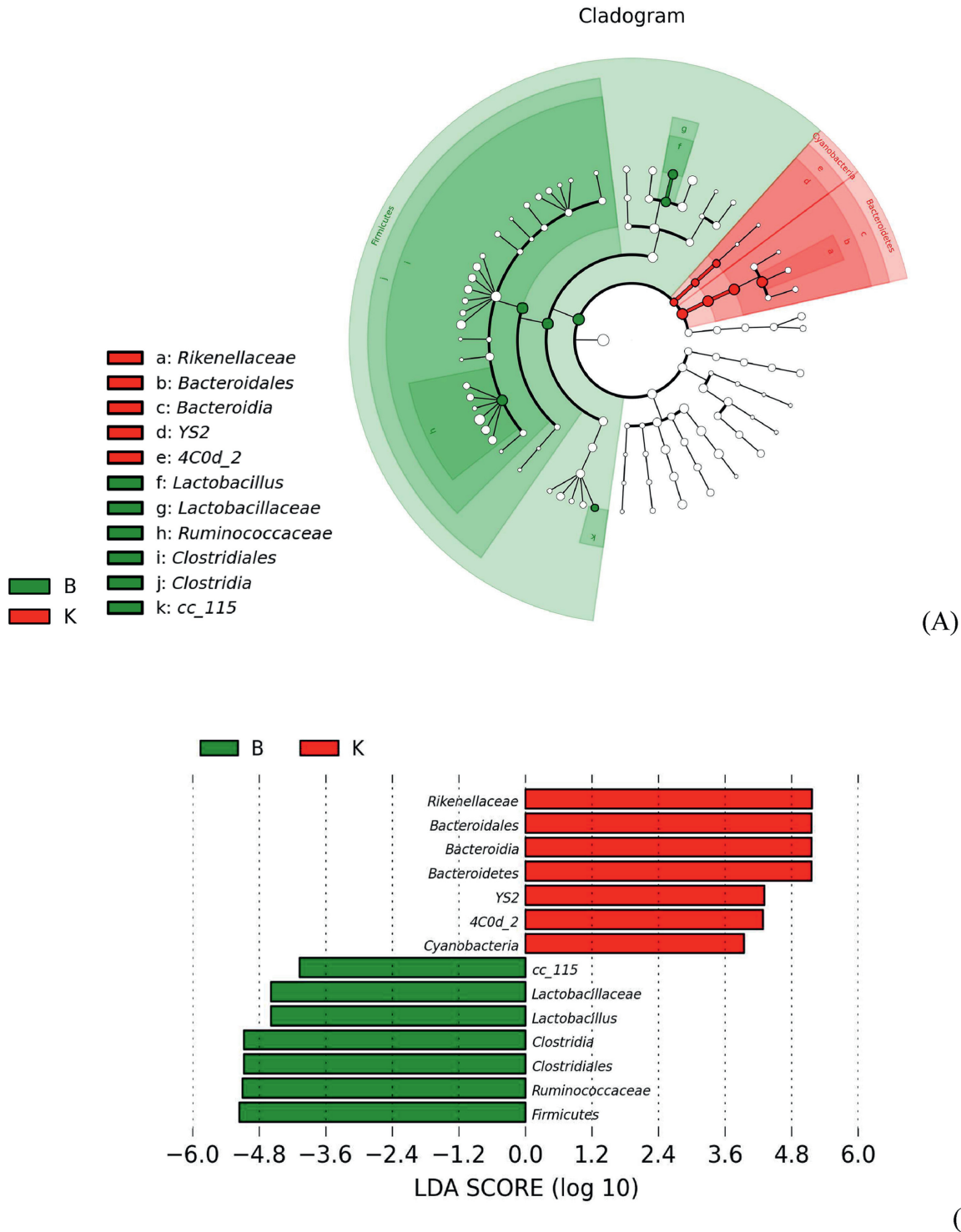
The gut microbiota play an important role in host health, disease, feed digestion and nutrient absorption, and production of poultry (Waite and Taylor, 2015). This study revealed that *E. coli* challenge dramatically altered alpha diversity of the cecal microbial community, while the changes in cecal bacterial diversity induced by *E. coli* O78 challenge were reversed by EOA addition, suggesting that EOA supplementation could prevent intestinal microbial disturbances caused by *E. coli* O78 challenge. In contrast to our results, several previous studies have reported that alpha



**Fig. 5.** Stacked bar chart of cecal microbial structure at the genus level. (A) Relative abundance of top 21 bacterial genera. (B) Comparison of the relative abundances of the bacterial genera which were greater than 1% among the 4 groups. Treatment groups: A = neither infection nor essential oil and organic acid mixture (EOA) addition control, K = *Escherichia coli* O78 infection control, B = EOA addition but without infection, L = both EOA addition and *E. coli* O78 infection.

diversity was unaffected by essential oils (Zhu et al., 2019) or organic acids (Bortoluzzi et al., 2017) or an EOA mixture (Pham et al., 2020, 2021). The variability in results was possibly associated with different EOA products and challenge models. There could be a possibility that the EOA treatment effectively reduced the “bad” bacteria and promoted the “good” bacteria when broiler chickens were confronted with the pathogen challenge, but this hypothesis needs to be further evaluated. PLS-DA and NMDS analysis showed that EOA addition, *E. coli* O78 challenge, and both

changed the beta diversity of the cecal microbial community, indicating that these treatments significantly affected the intestinal bacterial community profile. Furthermore, LEfSe analysis showed that EOA administration could improve cecal microbial composition in unchallenged birds, as evidenced by increased abundance of some beneficial bacteria, such as Ruminococcaceae, Lactobacilaceae and *Lactobacillus*, which was in accordance with several previous findings (Yin et al., 2017; Yang et al., 2019; Pham et al., 2020, 2021), indicating that EOA supplementation was good

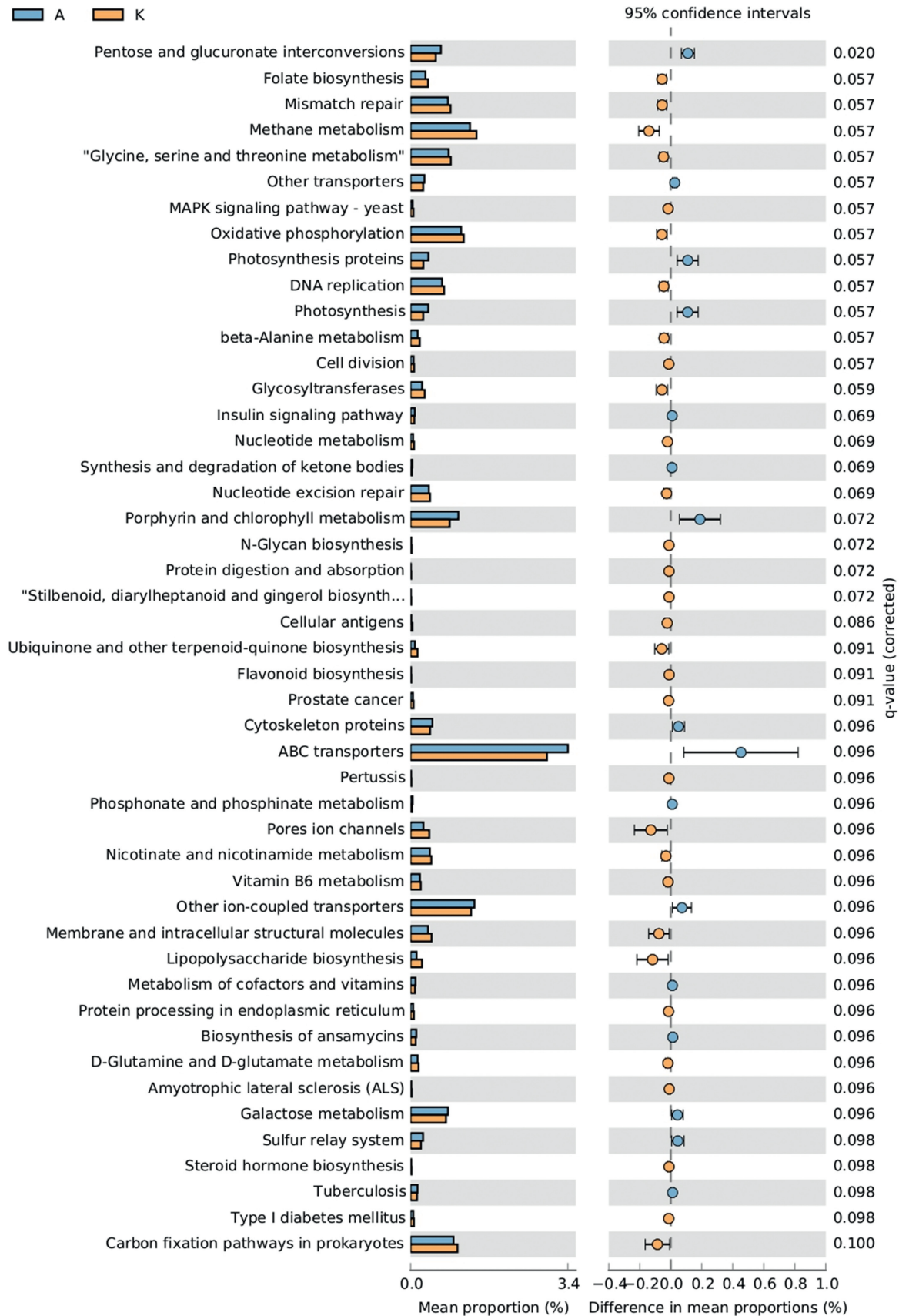


**Fig. 6.** Differential bacterial taxa in cecal microbiota of the B and K groups using LEfSe analysis ( $P < 0.05$  and LDA cutoff  $> 2.0$ ) ( $n = 6/\text{group}$ ). (A) Cladogram plot. (B) The histogram shows differentially abundant bacteria in the B and K groups ranked by linear discriminant analysis (LDA) scores. Treatment groups: A = neither infection nor essential oil and organic acid mixture (EOA) addition control, K = *Escherichia coli* O78 infection control, B = EOA addition but without infection, L = both EOA addition and *E. coli* O78 infection.

for gut health of broiler chickens reared under conditions without systemic infection.

Interestingly, in our study, *E. coli* intratracheal infection alone was observed to remarkably decrease the relative population of Firmicutes, but enrich the percentage of *Lactobacillus* and *Oscillospira* compared to the non-challenged control, while data from

the infected birds fed EOA appeared to decrease cecal *Bacteroidetes* and *Lactobacillus* abundance compared with the single infected control, which was opposite to what was observed in the *Lactobacillus* population in the non-challenged broiler chickens supplemented with EOA. In the cecal microbiota, different trends in relative abundance of total *Lactobacillus* and different strains of



**Fig. 7.** Predictive functional profiles generated from 16S rRNA marker gene sequences using PICRUSt at KEGG levels 2. (A and B) Differentially regulated metabolic pathways in A vs. K group, K vs. L group, respectively. The corrected *P*-value is listed at the right. Treatment groups: A = neither infection nor essential oil and organic acid mixture (EOA) addition control, K = *Escherichia coli* O78 infection control, L = both EOA addition and *E.coli* O78 infection.

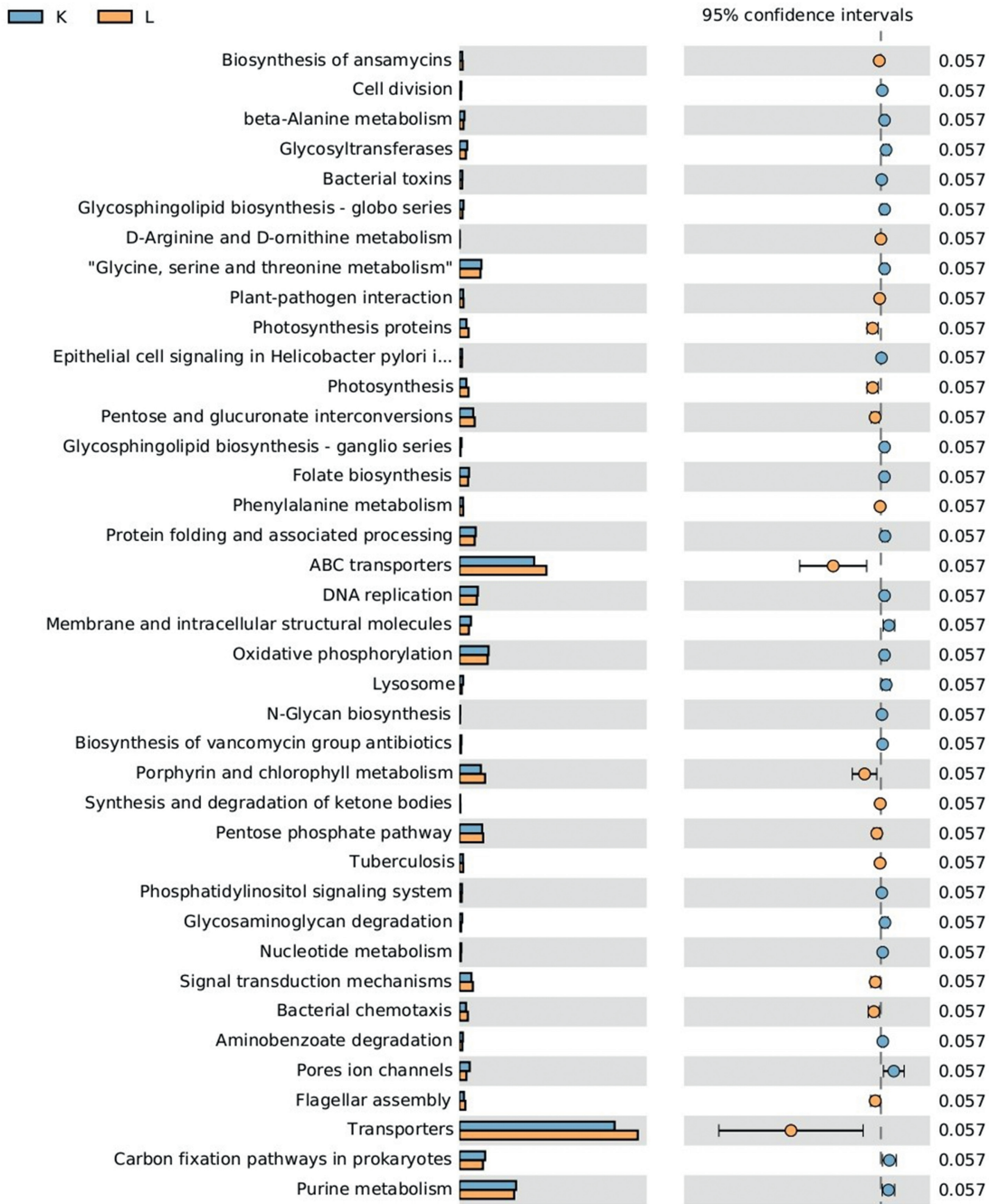


Fig. 7. (continued).

*Lactobacillus* of cecal microbiota were been observed in non-challenged or challenged broiler chickens given EO, OA, or EOA from previous studies. Some authors have suggested that supplemental EO, OA or EOA could increase the proportion of *Lactobacillus* spp. in chickens without challenge (Abudabos and Al-Mufarrej, 2014; Yang et al., 2019; Dai et al., 2021). Other researchers found that with the appropriate dosage, EO addition could increase *Lactobacillus* populations, whereas a high dosage of EO inhibited the growth of *Lactobacillus* in birds challenged with *C. perfringens*, and this study also indicated that EO administration differentially modulated the abundance of different strains of *Lactobacillus* in challenged birds (Du et al., 2015; Yin et al., 2017). Conversely, several authors reported that supplementation with EO decreased

cecal *Lactobacillus* abundance in the unchallenged broiler chickens (Zhu et al., 2019; Chen et al., 2020). Discrepancies in the reported results might be due to one or several factors, including the chemical composition of EO, OA, and EOA blend coating, challenged or not, challenge model, as well as time of sampling post challenge. *Lactobacillus* is the most abundant beneficial bacteria in the gut, which confers anti-inflammatory responses and affects gut integrity via modulating mucosal immune responses (Bermudez-Brito et al., 2012; Ashraf and Shah, 2014). Numerous studies have suggested that *Oscillospira* play an important role in gut health and metabolic diseases, and is both positively and negatively associated with many diseases (Konikoff and Gophna, 2016; Yang et al., 2021). *Bacteroides* species, the gut's most common anaerobes, have been

documented to create lipopolysaccharides (LPS), impact host immunity, and inhibit pathogen colonization of the gastrointestinal tract (GIT) (Hiippala et al., 2018; Zafar and Saier, 2021). A higher proportion of *Lactobacillus* and *Oscillospira* was observed in the cecal contents of the single *E. coli* O78-challenged birds, suggesting that *E. coli* O78 infection in fact promotes the growth of intestinal beneficial bacteria, resulting in an enhanced host anti-inflammatory response. However, a lower percentage of cecal *Bacteroidetes* and *Lactobacillus* which was discovered in APEC-infected broiler chickens after they were fed EOA, may be linked to the restoration of intestinal microbiota balance, reduced gut inflammation, and fewer gut lesions. As a result, the findings of this study found that EOA improved FCR and gut health in APEC-infected broiler chickens, as well as reducing gut inflammation, most likely due to favorable modification of intestinal microbiota composition. However, the correlation among gut microbiota, APEC challenge, EOA treatment and poultry gut health requires further investigation.

Most surprisingly, PICRUSt analysis revealed that only the pentose and glucuronate interconversion pathway of cecal microbiota was suppressed after *E. coli* O78 infection, while EOA addition had no significant influence on the function of cecal microbiota of broiler chickens regardless of *E. coli* O78 challenge. The pentose and glucuronate interconversion pathway have been reported to be involved in sugar utilization in microorganisms and redox status and therefore could provide nucleic acid synthesis substrates and keeps glutathione in its reducing state (Lakshmanan et al., 2021). Thus, the pentose and glucuronate interconversion pathway of cecal microbiota was suppressed after *E. coli* O78 infection, suggesting that *E. coli* O78 infection might cause oxidative stress in the gut through downregulating carbohydrate-related metabolic pathways. Whether the supplementation of EOA improves gut health through its antioxidant pathway requires further investigation. In addition, the impacts of EOA on fecal metabolite profiles should be investigated in order to understand the causal relationships between EOA, metabolites, and intestinal function.

## 5. Conclusion

The results of this study suggested that supplemental coated essential oil and organic acid mixture remarkably improved feed conversion efficiency, lowered gross lesion scores and cecal *E. coli* population, altered cecal microbiota composition, inhibited mucosal and systemic inflammation, enhanced intestinal goblet cells and serum IgG concentration along with ZO-1 gene expression of the infected birds compared with the infected groups alone. Together, the coated essential oil and organic acid mixture could be used as an antibiotic alternative to alleviate *E. coli* O78-induced gut injury and inflammation in the broiler industry.

## Author contributions

**Van Hieu Pham:** Investigation, Conceptualization, Data curation, Writing - original draft, Validation. **Jinyu Huang:** Conceptualization, Methodology. **Fangshen Guo:** Conceptualization, Methodology, Writing - review & editing. **Kaichen Zhang:** Visualization, Investigation. **Linhua Kong:** Investigation. **Wenrui Zhen:** Methodology, Software. **Waseem Abbas:** Writing - review & editing. **Zhong Wang:** Conceptualization, Methodology, reviewing & editing, Supervision. **Yuming Guo:** Reviewing & editing, Supervision, Project administration.

## Declaration of competing interest

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

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