Dietary supplementation with fermented defatted “alperujo” induces modifications of the intestinal mucosa and cecal microbiota of broiler chickens

Agustín Rebollada-Merino,* María Ugarte-Ruiz,*,1 Marta Hernández,†,‡ Pedro Miguela-Villoldo,*,# David Abad,† Pedro Cuesta-Alvaro,*, David Rodríguez-Lázaro,† Lucía de Juan,*,‡ Lucas Domínguez,*,‡ and Antonio Rodríguez-Bertos*†‡

*VISA VET Health Surveillance Centre, Complutense University of Madrid, 28040 Madrid, Spain; †Laboratorio de Biología Molecular y Microbiología, Instituto Tecnológico Agrario de Castilla y León, 47071 Valladolid, Spain; ‡Área de Microbiología, Departamento de Biotecnología y Ciencia de los Alimentos, Universidad de Burgos, 09001 Burgos, Spain; #Department of Animal Health, Faculty of Veterinary Medicine, Complutense University of Madrid, 28040 Madrid, Spain; and †‡Department of Internal Medicine and Animal Surgery, Faculty of Veterinary Medicine, Complutense University of Madrid, 28040 Madrid, Spain

ABSTRACT Nutraceuticals are not only nutritionally beneficial for animals but also their use as feed supplements may reduce environmental contamination. The effect of fermented defatted “alperujo,” an olive oil by-product, supplementation on the intestinal health of broiler chickens was assessed by analyzing the intestinal mucosal morphology of the duodenum and the cecum. The microbiota of the cecum was also characterized by analyzing the V3-V4 region of the 16S rRNA gene on days 7, 14, 21, 28, 35, and 42. Supplemented broilers from 14 to 35 D of age showed an increase in villus height in the duodenum. This increase likely improved digestibility and absorption capacity during growth, leading to the observed increase in BW at day 35 of life. A progressive increase in crypt depth in both the duodenum and the cecum was also observed. This modification likely enhanced epithelial renewal, thus safeguarding the turnover capacity of the intestinal mucosa. Our molecular analysis of cecal microbiota suggests that this dietary supplement may favor the growth of certain bacteria and may control the spread of pathogenic bacteria by means of competitive exclusion.

Key words: intestinal health, olive oil by-product, fermented defatted “alperujo”, histology, microbiota

INTRODUCTION

A healthy gut is critical for the general health and welfare of poultry (Ducatelle et al., 2018). Intestinal health is a permanent balance that is influenced by the interactions between the gastrointestinal mucosa and environmental factors such as diet and infectious agents, which in turn, affect the microbiota of the gut (Yegani et al., 2008). Consequently, the mucosa and microbiota are the main components involved in determining intestinal health, as well as modulating the immune response (Yegani and Korver, 2008; Lee et al., 2017; Ducatelle et al., 2018).

The intestinal mucosa is a dynamic organ that undergoes continuous renewal. Damage to the epithelial surface leads to morphologic changes in the structure of villi and crypts (Ducatelle et al., 2018), thus interfering with the proper functioning of the intestinal mucosa. Epithelial injuries result in deficiencies in nutrient and water absorption and cause an unspecific immune response. A diet-based approach that establishes an adequate immune response during early production phases may prevent infectious diseases, a common cause of production losses in the poultry industry (Yegani and Korver, 2008; Rubio, 2019). On the other hand, the intestinal microbiota, and its interactions with the mucosa and other environmental factors, plays an important role in the intestinal development and physiology of poultry. Its role in digestion,
absorption, and metabolism could directly influence immunity and performance (Yadav and Jha, 2019).

In general, diet composition is the main factor that influences the early development and health of the gastrointestinal tract (Rubio, 2019). Differential intake of components of poultry feed produces changes in both mucosal morphology and microbiota (Ducatelle et al., 2018) and can even induce the selection and growth of certain bacterial strains (Kelly et al., 2017; Lee et al., 2017; Yadav and Jha, 2019). Numerous dietary supplements, particularly those considered probiotic, prebiotic, or nutraceutical, have been added to broiler chicken feed.

Nutraceuticals are substances derived from natural sources that have benefits other than for nutrition (Nasri et al., 2014). In recent years, a main focus of animal research, particularly of poultry science, has been on the ability of natural compounds to promote a healthy gut by modifying mucosal morphology and microbiota, thereby improving the overall health status of chickens. Supplementation with natural compounds has improved the productive performance of some farms (Rubio, 2019). Some natural compounds, which are primarily obtained from the food industry, may not only be nutritionally valuable for animals but also their use as feed supplements may help reduce environmental contamination (Viveros et al., 2011; Berbel and Posadillo, 2018; Pappas et al., 2019). In particular, olive oil by-products, given their high polyphenolic content, have been associated with antioxidant, anti-inflammatory, antithrombotic, antimicrobial, and anti-coccidian effects in poultry (Alu’datt et al., 2010; Gerasopoulos et al., 2015; Nasopoulou et al., 2018).

The two main centrifugation systems used to obtain olive oil generate different types of wastes (Alburquerque et al., 2004). The three-phase extraction system generates olive mill wastewater and olive cake or pulp, which, as dietary supplements, have been shown to improve the organic antioxidant capacity, health status, and productive performance of broilers (Gerasopoulos et al., 2015; Sabino et al., 2018; Papadomichelakis et al., 2019). However, since the early 1990s, two-phase centrifugation has been the most implemented system, at least in the Spanish olive oil industry (Alburquerque et al., 2004). The two-phase system generates a waste product referred to as two-phase mill waste, “alperujo” or olive pomace. Of all the olive oil by-products, this one has been one of the most common to be tested as a supplement in animal feed (Berbel and Posadillo, 2018). Olive pomace extract has been reported, in particular, to have anti-inflammatory properties that enhance gut function in broilers (Herrero-Encinas et al., 2020).

Despite its beneficial effects and in contrast to other food industry wastes, further processing of “alperujo” is required before it can be used in animal feed (Berbel and Posadillo, 2018). “Alperujo” that had been previously fermented and defatted was recently shown to enhance intestinal health in laying hens by modifying the intestinal mucosa and microbiota, in addition to improving shell hardness (Rebollada-Merino et al., 2019). After degreasing, the free phenols present in “alperujo” appear to have the same level of antioxidant activities as those in full-fat olive cake (Alu’datt et al., 2010). This observation suggests that modified olive oil by-products, which can be used directly as dietary supplements, possess the same properties as unmodified ones. However, the effect of fermented defatted “alperujo” on broiler intestinal health has not been studied to date.

To evaluate the effect of feed supplementation on intestinal health, a multidisciplinary approach is required. In this context, histomorphologic and microbiota analyses have been shown to be reliable biomarkers in chickens (Viveros et al., 2011; Ducatelle et al., 2018). The aim of this experimental study was to analyze the potential effect of the modified olive oil by-product fermented defatted “alperujo” on the intestinal health of broilers by characterizing changes in the mucosal morphology of the duodenum and the cecum and in the microbiota of the cecum throughout the production cycle.

MATERIALS AND METHODS

Ethical Approval

Experimental procedures were approved by the Complutense University of Madrid Animal Care and Ethics Committee in compliance with the Community of Madrid (PROEX 152/19). Research met the guidelines approved by the Institutional Animal Care and Use Committee.

Birds, Rearing Conditions, and Diet

Sixty 1-day-old male Ross 308 broiler chickens were obtained from a commercial hatchery and raised in the laboratory facilities of the VISAVET Health Surveillance Center for 42 D under the same environmental and light conditions previously used for the species (Herrero-Encinas et al., 2020). Chickens were physically separated on arrival into 2 groups: a control group fed with conventional feed and a treated group whose feed was supplemented with 2% fermented defatted “alperujo” that was provided by Porres y Barios, S.A. (Córdoba, Spain) and the composition and processing to obtain fermented defatted “alperujo” was previously described by Rebollada-Merino et al. (2019). Animals had ad libitum access to feed and water for the duration of the experiment. They were monitored physically twice daily and at all times by video cameras.

Postmortem Study and Sample Collection

On posthatching days 7, 14, 21, 28, 35, and 42, five randomly selected animals from each group were sedated intramuscularly with diazepam and euthanized with an overdose of sodium pentobarbital by intravenous injection. Before the necropsy, each animal was weighed. A completed and systematic postmortem was performed,
and duodenum and cecum samples were collected and fixed in a 10% formaldehyde-buffered solution (Panreac Química SLU, Barcelona, Spain). In addition, the cecal content was collected from each animal and preserved at −80°C for metagenomics studies.

**Histomorphometric Study**

After routine histopathologic processing and hematoxylin–eosin staining (Rebollada-Merino et al., 2019), sections of the duodenum and the cecum were examined under an optical microscope coupled with a digital camera, and a blind histomorphometric analysis was performed at 40 × magnification. Using an image analyzer (Leica Application Suite; Leica, Wetzlar, Germany), 20 intact and well-oriented villi and 20 crypts in the duodenum, as well as 20 crypts in the cecum, were evaluated per animal.

**Statistical Analysis**

Differences between the two groups by age were assessed using the Mann-Whitney test implemented in IBM SPSS Statistics v25 (IBM, Armonk, NY). Statistical significance was considered at $P < 0.05$.

**Total DNA Extraction and 16S rRNA Library Preparation and Sequencing**

Total DNA was extracted from 220 mg of the cecal content using the QIAamp DNA Stool Mini Kit (Qiagen, Hilden, Germany), as per the manufacturer’s instructions. DNA concentration was determined using a Qubit fluorometer (Invitrogen, Carlsbad, CA). Microbial diversity was studied by analyzing sequences of the V3-V4 region of the 16S rRNA gene. The primers and PCR conditions used for this analysis were as previously reported (Klindworth et al., 2013). Sample multiplexing, library purification, and sequencing were carried out as described in the “16S Metagenomic Sequencing Library Preparation” guide by Illumina (San Diego, CA). Libraries were sequenced on an Illumina MiSeq platform that provided 300-bp paired-end reads.

**Bioinformatics and Data Analysis**

Raw demultiplexed sequence data were processed using QiimeReporter (https://github.com/dabadgarcia/qiime_reporter). This straightforward pipeline for the analysis of amplicon sequences integrates basic Qiime2 commands (Bolyen et al., 2019) with R programming language. In brief, the DADA2 package (Callahan et al., 2016) was used to filter reads, merge paired ends, remove chimeras, and assign amplicon sequence variants. Then, a pretrained Naïve Bayes classifier (Wang et al., 2007) was used to obtain the taxonomic assignment of the amplicon sequence variants, using the SILVA database v132 (Quast et al., 2013) as a reference, which resulted in a table containing the microbial composition for each of the samples. Raw reads are available in the BioProject database with ID PRJNA643396 (http://www.ncbi.nlm.nih.gov/bioproject/643396).

**RESULTS**

**Body Weight**

On posthatching days 7, 14, 21, and 28, no BW differences were observed between the control and supplemented groups (Figure 1). However, at 35 D, the mean BW of the supplemented group was significantly higher than that of the control ($P = 0.009$). At 42 D, the mean weight of the supplemented group was still higher, although the difference was not statistically significant ($P = 0.117$).

**Histomorphometric Study**

The histomorphometric analysis of the duodenum revealed a significant increase ($P < 0.05$) in villus height on days 14, 21, 28, and 35 in the fermented defatted “alperujo”–supplemented group (Table 1). Duodenal crypts were significantly deeper in the control group on days 14 and 42 ($P < 0.05$); however, the mean values, although not significant ($P > 0.05$), were higher in the supplemented group for the other days. In the cecum, crypts were significantly deeper in the supplemented group for all samplings ($P < 0.05$), except on day 14 (Table 1, Figure 2).

**Cecal Microbiota**

Nonsignificant variability in bacterial taxonomic richness at the genus level was observed in the treated animals (Figure 3). The most abundant bacterial genera identified at day 7 were *Clostridiales* (relative abundance: 30.15%), *Klebsiella* (29.27%), and *Escherichia-Shigella* (22.99%) in the control group and *Escherichia-Shigella* (32.06%), *Ruminococcus* (25.63%), and *Lachnospiraceae* (16.22%) in the treated group. At day 14, *Escherichia-Shigella* (45.30 and 39.68% in control and treated, respectively) and *Lachnospiraceae* (20.21 and 9.71%) were most prevalent in both groups, though in the treated group, *Oscillibacter* (34.62%) was also highly abundant. At day 21, *Escherichia-Shigella* (48.78%) and *Lactobacillus* (11.45%) were the most abundant genera in the control group. By contrast, in the treated group, *Escherichia-Shigella* (22.11%), *Oscillibacter* (20.47%), and *Lachnospiraceae* (16.65%) were most abundant at this stage. At day 28, *Lachnospiraceae* (19.41 and 21.49% in control and treated, respectively) were most abundant in both groups. At day 35, *Bacteroides* (24.16%) and *Lachnospiraceae* (23.51%) were most abundant in the control and treated groups, respectively, followed by *Escherichia-Shigella* in both groups (13.21% in the control and 15.77% in the treated). Finally, at day 42, *Bacteroides* was the most abundant genus in both groups (35.87% in control and
53.52% in the treated), followed by Lachnospiraceae in the control group (15.64%) and *Escherichia-Shigella* in the treated group (8.13%).

**DISCUSSION**

Diet supplementation with nutraceuticals, natural compounds with positive health effects, is an active field of research in veterinary medicine, particularly in poultry science. Dietary changes have been shown to contribute to structural and bacterial modifications in the intestine that improve intestinal health and, consequently, poultry production performance (Viveros et al., 2011; Rebollada-Merino et al., 2019; Yadav and Jha, 2019). After fermented defatted *alperujo* supplementation, we observed a significant increase in BW at day 35 of life, which may be confirmed in further experiments with a higher sample size. Tufarelli et al. (2016) previously reported a BW increase after olive oil supplementation in broilers and suggested that lipid content directly influences weight gain and feed efficiency. Increased BW has also been attributed to polyphenols present in olive oil and olive oil by-products (Tufarelli et al., 2015). However, the results of our histomorphometric study of the duodenum and cecum suggest improved nutrient absorption in the intestine may be associated with morphologic changes in the mucosa in addition to the nutrient composition of the supplement. Histologic measurements are considered the gold standard for evaluating intestinal health (Samuel et al., 2017) and the essential oil carvacrol (Kelly et al., 2017). In addition, supplementation with *Pulicaria gnaphalodes* powder containing a high phenol content has been associated with a reduction of pathogenic bacteria in the small intestine, which may be due to increased mucus secretion by goblet cells (Shirani et al., 2019). Increases in villus height are thought to be caused by the action of more transport enzymes in the enterocytes (Viveros et al., 2011; Ma et al., 2018).

We observed a significant increase in duodenal villus height in supplemented animals from the second to fifth week of life. This increase may result in a greater absorption capacity, which may partially explain the greater weight increase observed in this group, especially in the final week of production (Viveros et al., 2011; Samuel et al., 2017; Reis et al., 2018). A similar increase in villus height has been reported in broilers supplemented with the plant-derived phenol gallic acid (Samuel et al., 2017) and the essential oil carvacrol (Kelly et al., 2017). In addition, supplementation with *Pulicaria gnaphalodes* powder containing a high phenol content has been associated with a reduction of pathogenic bacteria in the small intestine, which may be due to increased mucus secretion by goblet cells (Shirani et al., 2019). Increases in villus height are thought to be caused by the action of more transport enzymes in the enterocytes (Viveros et al., 2011; Ma et al., 2018). However, a recent study has suggested that phenolic compounds inhibit digestive enzymes and transporters

---

**Table 1.** Results of the histomorphometric analysis of the duodenum (villus height and crypt depth) and cecum of control (*n* = 100) and supplemented (*n* = 100) broilers. Number of samples (N), mean values in micrometers (Mean), SD, and P-value are shown for each age group.

|                  | Control | Supplemented | P-Value<sup>1</sup> |
|------------------|---------|--------------|---------------------|
| **Duodenal villus height** |         |              |                     |
| 7 D              | 702.31  | 670.70       | 236.95              |
| 14 D             | 1,068.80| 1,168.35     | 177.21              |
| 21 D             | 857.38  | 1,303.24     | 441.13              |
| 28 D             | 1,003.97| 1,422.50     | 264.57              |
| 35 D             | 990.49  | 1,199.55     | 286.47              |
| 42 D             | 1,196.92| 1,103.85     | 339.67              |
| **Duodenal crypt depth** |         |              |                     |
| 7 D              | 99.13   | 105.41       | 46.84               |
| 14 D             | 137.32  | 123.99       | 48.23               |
| 21 D             | 170.49  | 158.08       | 59.72               |
| 28 D             | 131.40  | 122.11       | 88.47               |
| 35 D             | 113.35  | 108.03       | 132.68              |
| 42 D             | 143.47  | 130.43       | 132.68              |
| **Cecal crypt depth** |         |              |                     |
| 7 D              | 139.44  | 207.81       | 86.84               |
| 14 D             | 217.02  | 207.60       | 59.38               |
| 21 D             | 295.57  | 449.94       | 175.81              |
| 28 D             | 391.66  | 492.70       | 245.29              |
| 35 D             | 294.92  | 370.86       | 122.90              |
| 42 D             | 276.64  | 617.00       | 236.68              |

<sup>1</sup>The Mann-Whitney test was used to assess significant differences between control and supplemented animals.

---

**Figure 1.** Effect of supplementation with 2% fermented defatted “alperujo” on broiler BW (in g) by age. At day 35, the mean BW was significantly higher in the treated group than that in the control group (*P* = 0.009).

**Figure 2.** Cecum of 28-day-old broilers. Hematoxylin–eosin stained section of a control (A) and a supplemented (B) cecum. Crypts were significantly deeper in the supplemented group than those in the control group. 40×, scale bar: 500 μm.
in the enterocytes, delaying the digestion of some macro-
nutrients, such as proteins and carbohydrates, and even
hindering fat absorption (Domínguez-Avila et al., 2017).
Our results indicate that villus growth in the duodenum
of supplemented broilers is dynamic but not linear: a
progressive increase in intestinal villus height was
observed until it reached a maximum during the fourth
week of life, after which it markedly decreased. Such dy-
namics have not been previously reported in broilers,
mainly because other morphologic studies mainly focus
on the last phases of the production cycle.
In chicken intestines, epithelial cells in the villi are
continuously renewed by crypt stem cells (Ducatelle
et al., 2018). In broilers, the presence of deep crypts
generally implies a better response to potentially harm-
ful events in the villi (Pourabedin and Zhao, 2015;
Ducatelle et al., 2018; Sabino et al., 2018). In our study,
crypts, particularly in the cecum, were deeper in the
treated animals. By contrast, supplementation with
grape products or olive oil wastewater (a three-phase
mill waste), both of which contain high levels of polyphen-
ols, decreases crypt depth in broilers (Viveros et al.,
2011; Sabino et al., 2018). Compared with our study,
in the olive oil wastewater study by Sabino et al.
(2018), animals were supplemented with a lower by-
product concentration for less time, and fewer samplings
were taken for histologic analysis. Therefore, methodo-
logical differences may, in part, account for the contrast-
ing observations between studies. Regardless, in our
study, greater progressive growth in crypt depth was
observed until the fifth week of life, suggesting that fer-
mmented defatted “alperujo” improves epithelial renewal,
thus safeguarding the turnover capacity of crypts, which
is critical in the response to potential mucosal injuries,
secondary bacterial infections, or parasitic diseases.
Metagenomic studies of changes in cecal microbiota
can provide insight into potential intestinal health ben-
efits and host–microbiota interactions, despite only
providing limited information on the effect of bioactive
compounds on microbiota (Herrero-Encinas et al.,
2020; Yadav and Jha, 2019; Rychlik, 2020). As physio-
logical changes in intestinal microbiota have been associ-
ated with factors such as age and feed composition
(Videnska et al., 2014; Lee et al., 2017; Ijaz et al.,
2018; Richards et al., 2019; Rychlik, 2020), we evaluated
the microbial composition of the cecum, a main site of
bacterial metabolic activity (Cressman et al., 2010;
Richards et al., 2019; Yadav and Jha, 2019; Rychlik,
2020), throughout the productive life of broilers.
In our microbiota analysis, Enterobacteriaceae was
the most prominent family, consistent with the findings
of a previous study analyzing cecal microbiota in broilers

![Figure 3. Bar chart showing the 11 most abundant bacterial groups found in the cecal content of control and treated broiler chickens. Each bar represents the relative abundance of bacteria by animal, diet treatment, and age. Relative abundance is shown on the vertical axis and age (7, 14, 21, 35, or 42) and diet (C, control; T, treatment) on the horizontal axis.](image-url)
after *Lactobacillus acidophilus* supplementation (De Cesare et al., 2017). Specifically, during the first day of life, *Escherichia-Shigella* and *Klebsiella* were the most represented genera; however, their relative abundances decreased progressively thereafter. These results are in line with those reported by Ijaz et al. (2018) and Rychlik (2020). Some authors have hypothesized that a high level of dietary fiber accounts for the reduction of these bacteria in the gut (Walugembe et al., 2015). The fermented defatted “alperujo” supplement also has a high fiber content (Rebollada-Merino et al., 2019), supporting the idea that fiber hinders the growth of Enterobacteriaceae. In addition, some Enterobacteriaceae genera may be pathogenic (Ma et al., 2018), thus fermented defatted “alperujo” supplementation in broilers may act to reduce the abundance of these bacteria in the gut, resulting in less epithelial damage and inflammation. The reduced abundance of these bacteria in the cecum potentially explains the observed differences in crypt depth between 21 and 28 D of life.

In contrast, *Oscillibacter* abundance increased in the supplemented group from 14 to 35 D of life. *Oscillibacter* is a butyrate-producing bacterium that is hypothesized to increase short-chain fatty acid availability in the intestinal lumen (Rychlik, 2020).

At 42 D, the microbiota of supplemented broilers was, in general, similar to that of the control. One difference was the higher relative abundance of *Bacteroides*, at the expense of Lachnospiraceae and *Ruminococcus*, in the supplemented group. *Bacteroides*, which has been reported to be predominant in broilers around this age (Lee et al., 2017), increases propionic acid generation in the cecum, which may correspond to a reduction of fat synthesis in the liver (Qi et al., 2019) that, in turn, prevents hepatic steatosis.

We also noted some contradictory results in our analysis of microbiota composition with respect to the abundance of Ruminococcaceae and *Lactobacillus* and overall bacterial diversity. Ruminococcaceae has been associated with an increase in BW after *Bacillus subtilis* supplementation in broilers (Ma et al., 2018). In our study, Ruminococcaceae was underrepresented in the supplemented animals, yet we found significant differences in the BW of 35-day-old broilers between the two groups. This result suggests that low levels of Ruminococcaceae do not necessarily decrease BW and that the BW increase observed in fermented defatted “alperujo”–supplemented animals is likely influenced by some other microbial interaction. *Lactobacillus* appears to reduce the antigen load from resident bacteria, thus having an anti-inflammatory effect in the gut (De Cesare et al., 2019). Here, we observed a lower relative abundance of *Lactobacillus* in the supplemented group at all time points, except 35 D of life, which is also when crypt depth diminished in these animals. Finally, in contrast with other broiler studies, we found one of the highest levels of bacterial diversity at 42 D of life (Herrero-Encinas et al., 2020; Kumar et al., 2019).

To summarize, fermented defatted “alperujo” supplementation in broilers improves villus height in the duodenum at early life stages. This likely increases digestibility and absorption capacity during growth, leading to an increase in BW at day 35 of life. Moreover, we suggest that the greater progressive increase of crypt depth observed in both the duodenum and the cecum after supplementation leads to enhanced epithelial renewal, thus safeguarding the turnover capacity of the intestinal mucosa. Finally, the microbiota composition of supplemented broilers is dynamic: Enterobacteriaceae dominates during the first ws of life but then is replaced by *Oscillibacter* and Lachnospiraceae followed by *Bacteroides* at the end of the production cycle. These microbiota changes suggest that the fermented defatted “alperujo” dietary supplement may favor the growth of certain bacteria and may control the spread of pathogenic bacteria by means of competitive exclusion.

**ACKNOWLEDGMENTS**

This study was conducted as part of the project “Promoting One Health in Europe through joint actions on foodborne zoonoses, antimicrobial resistance and emerging microbiological hazards”. This project received funding from the European Union Horizon 2020 research and innovation programme under Grant Agreement No 773830.

The authors acknowledge Luis Arrabal (Porres y Barrios, S.A.) for providing the fermented defatted “alperujo” and VISAVET Health Surveillance Centre personnel involved in the project. Specifically, we thank the Pathology and Forensic Veterinary Unit: M. C. Jiménez, S. Cruz, G. Torre, F. Mayoral, N. Porras, L. Barreno; the Foodborne Zoonoses and Antimicrobial Resistance Unit: M. García, E. Rivero, M. Maaoumi; and the Quality and Biosafety Unit: M. Mazariiegos, L. Delgado, D. Duque, P. Alcubilla. We also thank English editing services provided by M. Modrell.

Conflict of Interest Statement: The authors did not provide a conflict of interest statement.

**REFERENCES**

Alburquerque, J. A., J. González, D. García, and J. Cegarra, 2004. Agricultural characterisation of “alperujo”, a by-product of olive oil extraction. Bioresour. Technol. 91:195–200.

Aludatt, M. H., I. Alli, K. Ereifej, M. Alhamad, A. Rahman Al-Tawaha, and T. Rababah, 2010. Optimization, characterization and quantification of phenolic compounds in olive cake. Food Chem. 123:117–122.

Berbel, J., and A. Posadillo, 2018. Review and analysis of alternatives for the valorization of agro-industrial olive oil by-products. Sustainbility 10:237.

Bolyen, E., J. R. Rideout, M. R. Dillon, N. A. Bokulich, C. C. Abnet, G. A. Al-Ghalith, H. Alexander, E. J. Alm, M. Arumugam, F. Asnicar, Y. Bai, J. E. Bisanz, K. Bittinger, A. Brejnrod, C. J. Brislawn, C. T. Brown, B. J. Callahan, A. M. Caraballo-Rodríguez, J. Chase, E. K. Cope, R. Da Silva, C. Diener, P. C. Dorrestein, G. M. Douglas, D. M. Durall, C. Duvallet, C. F. Edwardson, M. Ernst, M. Estaki, J. Fouquier, J. M. Gauglitz, S. M. Gibbons, D. L. Gibson, A. González, K. Gorlick, J. Guo, B. Hillmann, S. Holmes, H. Holste, C. Huttonower, G. A. Huttley, S. Jaussen, A. K. Jarmusch, L. Jiang, B. D. Kaehler, K. B. Kang, C. R. Keepe, P. Keim, S. T. Kelley, D. Knights, I. Koester,
Videnska, P., K. Sedlar, M. Lukak, M. Faldynova, L. Gerzova, D. Cejkove, F. Sisak, and I. Rychlik. 2012. Succession and replacement of bacterial populations in the caecum of egg laying hens over their whole life. PLoS One 9:e115142.

Viveros, A., S. Chamorro, M. Pizarro, I. Arija, C. Centeno, and A. Brenes. 2011. Effects of dietary polyphenol-rich grape products on intestinal microflora and gut morphology in broiler chicks. Poult. Sci. 90:566–578.

Walugembe, M., J. C. F. Hsieh, N. J. Koszewski, S. J. Lamont, M. E. Persia, and M. F. Rothschild. 2015. Effects of dietary fiber on cecal short-chain fatty acid and cecal microbiota of broiler and laying-hen chicks. Poult. Sci. 94:2351–2359.

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

Yadav, S., and R. Jha. 2019. Strategies to modulate the intestinal microbiota and their effects on nutrient utilization, performance, and health of poultry. J. Anim. Sci. Biotechnol. 10:2.

Yegani, M., and D. R. Korver. 2008. Factors affecting intestinal health in poultry. Poult. Sci. 87:2052–2063.