

## Effect of Dietary Inclusion of Ginger (*Zingier Officinal*) and Garlic (*Allium Sativum*) Oil Mixture on the Growth Performance and Caecal Microbial Population of Broiler Chickens

Oluwafemi RA<sup>1</sup>, Halima Abdullahi<sup>1</sup> and Alagbe JO<sup>2\*</sup>

<sup>1</sup>Department of Animal Science, University of Abuja, Nigeria

<sup>2</sup>Department of Animal Nutrition and Biochemistry, Sumitra Research Institute, Gujarat, India.

Received August 10, 2021; Revised October 10, 2021; Accepted October 13, 2021

### ABSTRACT

The objective of the present study was to determine effect of dietary inclusion of (*Zingier Officinal*) and garlic (*Allium sativum*) oil mixture (GIGM) on the growth performance and caecal microbial population of broiler chickens. One hundred and fifty-one-day-old broiler chicks (Ross 308) were randomly allocated into 5 treatments with three replicates consisting of 10 birds each in a completely randomized design. Birds in treatment 1 (T1) was fed basal diet with 0 % inclusion of GIGM while T2, T3, T4 and T5 were given 0.1 %, 0.2 %, 0.3 % and 0.4 % respectively. Clean feed and water were offered ad libitum and all other management practices were strictly observed throughout the experiment which lasted for 56 days. Results obtained were used to determine weight gain (WG), average daily weight gain (ADWG), total feed intake (TFI), average daily feed intake (ADFI), feed conversion ratio (FCR) and microbial population of *E. coli*, *Salmonella spp* and *Lactobacillus spp*. ADWG, ADFI and FCR were significantly ( $P < 0.05$ ) influenced by the dietary inclusion of GIGM. ADWG were highest in T5 (47.80 g), T4 (45.75 g) and T3 (45.09 g), intermediate in T2 (39.59 g) and lowest in T1 (30.72 g). *Lactobacillus spp* increased as the level of dietary inclusion of GIGM increases ( $P < 0.05$ ). *E. coli* and *Salmonella spp* counts were significantly ( $P < 0.05$ ) different among the treatments. It was concluded that GIGM could be included in the diet of broilers up to 0.4 % without causing any deleterious effect on the performance and health of birds.

### INTRODUCTION

Consumer pressure for antibiotic free poultry products has led to increased research in the area of antibiotic alternatives, including essential oils. Essential oils (EOs) are plant-based medicine that perform multiple biological activities such as; antimicrobial, antioxidant, antiviral, anti-inflammatory, antifungal, antiviral and hepato-protective [1]. According to Adewale [2] and Musa [3], EOs are volatile oily liquids extracted from plant parts, such as flowers, buds, stems, seeds, leaves, twigs and root which are capable of producing a positive physiological function in the body of animals. All plant parts synthesize an extremely diverse range of chemical compounds (phytochemicals) which represent a great potential for the discovery and development of new pharmaceuticals [4]. Among the essential oil of high medicinal value are ginger (*Zingier officinal*) and garlic (*Allium sativum*).

Ginger (*Zingier officinal*) belongs to the family Zingiberaceae. Its essential oil had long served the purpose of being medically significant, as antifungal, antibacterial, anti-inflammatory analgesic and immunodulatory impacts due to the presence of minerals, vitamins, amino acids and

phytochemicals ( $\beta$ -bisabolene and zingiberene (major) other sesquiterpenes include zingiberol,  $\alpha$ -curcumene,  $\beta$ -sesquiphellandrene,  $\beta$ -sesquiphellandrol (cis and trans); numerous monoterpenhydrocarbons, alcohols and aldehydes) [5-7].

Garlic (*Allium sativum*) contains sulfur compounds including alliin, produced enzymatically from (diallyl ehisosulfinate), allylpropyl disulfide, diallyl disulfide, diallyl trisulfide, ajoene and vicnyldithiines (secondary products of alliin produced non-enzymatically from alliin); S-allylmercaptocystocysteine (ASSC) and S-

**Corresponding author:** Alagbe JO, Department of Animal Nutrition and Biochemistry, Sumitra Research Institute, Gujarat, India, E-mail: demsonfarms@yahoo.com

**Citation:** Oluwafemi RA, Abdullahi H & Alagbe JO. (2023) Effect of Dietary Inclusion of Ginger (*Zingier Officinal*) and Garlic (*Allium Sativum*) Oil Mixture on the Growth Performance and Caecal Microbial Population of Broiler Chickens. J Microbiol Microb Infect, 5(1): 152-156.

**Copyright:** ©2023 Oluwafemi RA, Abdullahi H & Alagbe JO. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

methymercaptocysteine (MSSC); terpenes include citra, geraniol, linalool alfa and beta-phellandrene [8,9]. Previously, garlic oil has been listed as GRAS (Generally Recognized as Safe) [10].

The efficacies of EOs are well documented both *in vivo* and *in vitro* [11]. According to Burt [12], combination of essential oils has a greater antibacterial effect than individual EOs alone. EOs have been demonstrated to positively impact growth performance, blood profile and gut health of animals [13]. However, there are inconsistencies in the results due to differences in the chemical composition of EOs, which are affected by plant age or part used, extraction or processing methods, geographical locations and anti-nutrients [1].

Therefore, this experiment was designed to evaluate the effects of dietary inclusion of ginger (*Zingier officinal*) and garlic (*Allium sativum*) oil mixture on the growth performance and caecal microbial population of broiler chickens.

**MATERIALS AND METHODS**

**Experimental Site**

This study was carried out at the Department of Animal Science, University of Abuja Teaching and Research Farm, Main Campus, along airport Road, Gwagwalada, Abuja, Nigeria. Gwagwalada is the headquarters of the Gwagwalada Area Council located between latitudes 8°57' and 8°55'N and longitude 7°05' and 7°06' E [14].

**Sourcing and extraction of oil**

Fresh samples of ginger and garlic rhizomes were purchase from a local market in Gwagwalada Abuja, Nigeria. The samples were sorted out of the bad ones, then washed and peel manually with a kitchen knife to remove the outer covering of the rhizomes. It was dried for 14 days, milled into powder using a laboratory blender (Panasonic: Model 07A-08C) and then stored in an air tight well label container for further analysis. The oil was extracted using soxhlet extraction procedure; 100g of the sample were placed in a reflux condenser which consists of a condenser and a round bottom flask. The solvent used is petroleum ether and adjusted to 65°C to reach a vaporization point before the filtrate was exposed to the atmosphere and the residual solvent was allowed to evaporate before extracting the oil. The extracted oil was mixed in ratio 1: 1 to obtain ginger and garlic oil mixture (GIGLM).

**Experimental Animals and their management**

One hundred and fifty-one-day old (Ross 308) broiler chicks with mixed sex were used for the experiment. The birds were purchased from a commercial hatchery in Ibadan, Oyo State, Nigeria and weighed on arrival on the farm to obtain their initial body weight and thereafter weekly. A deep litter housing system was used for the experiment. Pens were fumigated two weeks prior to the commencement of the

study, surroundings were cleaned and foot bath was made available to ensure strict biosecurity. Birds were divided to five treatments with 3 replicates of ten birds in a completely randomized design. Charcoal pots were used as source of heat and wood shavings serve as the litter material. Vaccines were administered according to the disease condition in the environment and all other management practices were strictly adhered to throughout the experiment which lasted for 56 days.

**Diet formulation**

Two basal diets were formulated at different stages of production to meet up with the requirements of birds according to NRC (1994) as presented in **Table 1**. Broiler starter's mash (1-28 days) and finishers mash (29-56 days). Birds in Treatment 1 (T1) was fed dietary inclusion of ginger and garlic oil (GIGLM) at 0 %, while T2, T3, T4 and T5 were fed 0.1 %, 0.2 %, 0.3 % and 0.4 % respectively.

**Table 1.** Composition of the experimental diets.

Ingredients	Starter phase (%)	Finisher phase (%)
Maize	50.00	55.00
Soya bean meal	22.50	19.00
Groundnut cake	15.00	12.00
Fishmeal (72 %)	2.00	2.00
Wheat offal	4.45	6.05
Bone meal	2.00	2.00
Limestone	3.00	3.00
Salt	0.25	0.35
*Premix	0.25	0.25
Methionine	0.30	0.25
Lysine	0.25	0.20
Total	100.0	100.0
Calculated analysis (% DM)		
Crude protein (%)	23.05	21.41
ME (Kcal/kg)	2991.4	3100.3
Ether extract (%)	3.93	3.89
Crude fiber (%)	3.67	4.50
Calcium (%)	1.75	1.91
Phosphorus (%)	0.61	0.84

\*Premix supplied per kg diet: - vit A, 13,000 I.U; vit E, 5mg; vit D3, 3000I.U, vit K, 3mg; vit B2, 5.5mg; Niacin, 25mg; vit B12, 16mg; chorine chloride, 120mg; Mn, 5.2mg; Zn, 25mg; Cu, 2.6g; folic acid, 2mg; Fe, 5g; pantothenic acid, 10mg; biotin, 30.5g; antioxidant, 56mg.

**MEASUREMENTS**

**Performance parameters**

Feed intake (g) was determined by subtracting feed left over from feed served, it was estimated for each of the replicate daily.

Weight gain (g) = final weight - initial weight

Feed to gain ratio = feed intake (g)/weight gain (g)

Average daily weight gain (ADWG)  

$$= \frac{\text{Final body weight} - \text{Initial body weight}}{\text{Total days of the experiment}}$$

Average total feed intake (ADFI)  

$$= \frac{\text{Feed intake}}{\text{Total days of the experiment}}$$

**Caecal microbial enumeration**

On the 56<sup>th</sup> day of the experiment, 6 birds were randomly selected per treatment for caecal microbial enumeration (*E. coli*, *Salmonella spp* and *Lactobacillus spp*). A 10-fold serial dilution method was used in which 1% peptone solution was mixed with caecal samples and poured unto agar plates (Model R4-02X, Punjab, India) and incubated at 37°C for 48 h. Visible colonies were enumerated using colony counter and the results were expressed as log<sub>10</sub> CFU/g of caecal digesta.

**Phytochemical analysis**

Phytochemical analysis of GIGM was carried out using standard methods described by Harborne [15], Odebiyi and Sofowora [16].

**Statistical analysis**

Data obtained were subjected to one -way analysis of variance (ANOVA) using SPSS (23.0) and significant means

**Table 3.** Growth performance of broiler chickens fed dietary inclusion of ginger (*Zingiber officinale*) and garlic (*Allium sativum*) (GIGM) oil.

Parameter	T1	T2	T3	T4	T5	SEM
IBW (g)	42.80	42.90	43.10	43.00	43.20	0.22
FBW (g)	1720.4 <sup>c</sup>	2260.1 <sup>b</sup>	2568.3 <sup>a</sup>	2605.1 <sup>a</sup>	2720.0 <sup>a</sup>	10.82
WG (g)	1677.6 <sup>c</sup>	2217.2 <sup>b</sup>	2525.2 <sup>a</sup>	2562.1 <sup>a</sup>	2677.0 <sup>a</sup>	9.44
ADG (g)	30.72 <sup>c</sup>	39.59 <sup>b</sup>	45.09 <sup>a</sup>	45.75 <sup>a</sup>	47.80 <sup>a</sup>	1.05
TFI	4565.8 <sup>a</sup>	4243.8 <sup>b</sup>	4230.9 <sup>b</sup>	4031.4 <sup>b</sup>	3880.3 <sup>c</sup>	12.23
ADFI	81.53 <sup>a</sup>	75.78 <sup>b</sup>	75.55 <sup>b</sup>	72.00 <sup>b</sup>	69.29 <sup>c</sup>	2.44
FCR	2.72 <sup>a</sup>	1.91 <sup>b</sup>	1.70 <sup>b</sup>	1.60 <sup>c</sup>	1.50 <sup>c</sup>	0.12

were separated using the software of the same package. Significant was declared if P ≤ 0.05.

**RESULTS**

**Phytochemical composition of ginger (*Zingiber officinale*) and garlic (*Allium sativum*) oil**

The phytochemical composition of ginger (*Zingiber officinale*) and garlic (*Allium sativum*) oil (GIGM) is presented in **Table 2**. The sample contains flavonoids, saponins, terpenoids, phenols, oxalates, alkaloids and tannins at 20.78 %, 6.10 %, and 12.71 %, 17.90 %, 2.04 %, 10.31 % and 9.44 % respectively. In order of abundance flavonoids > phenols > terpenoids > alkaloids > tannins > saponins > oxalates.

**Table 2.** Phytochemical composition of ginger (*Zingiber officinale*) and garlic (*Allium sativum*) oil.

Constituents	Composition (%)
Flavonoids	20.78
Saponins	6.10
Terpenoids	12.71
Phenols	17.90
Oxalates	2.04
Alkaloids	10.31
Tannins	9.44

**Growth performance of broiler chickens fed dietary inclusion of ginger (*Zingiber officinale*) and garlic (*Allium sativum*) (GIGM) oil**

Growth performance of broiler chickens fed dietary inclusion of ginger (*Zingiber officinale*) and garlic (*Allium sativum*) oil is presented in **Table 3**. Initial body weight (IBW) ranged from 42.80 - 43.10 g, final body weight (1720.4 - 2720.0 g), weight gain (1677.6 - 2677.0 g), average daily weight gain (30.72 - 47.80 g), total feed intake (3880.3 - 4565.8 g), average daily feed intake (69.29 - 81.53 g) and feed conversion ratio (1.50 - 2.72). WG, ADG, TFI and ADFI were highest in T3, T4 and T5 relative to the other treatments (P<0.05). FCR were significantly different among the treatments (P<0.05).

Means in the same row with different superscripts differ significantly ( $P < 0.05$ ); T1: 0 % GIGM; T2: 0.1 % GIGM; T3: 0.2 % GIGM; T4: 0.3 % GIGM; T5: 0.4 % GIGM; IBW: initial body weight; FBW: final body weight; WG: weight gain; ADG: average daily weight gain; TFI: total feed intake; ADFI: average daily feed intake; FCR: feed conversion ratio.

**Caecal microbial population of broiler chicks fed diets with different level of GIGM**

Caecal microbial population of broiler chicks fed diets with different level of GIGM is presented in **Table 4**. Microbial population of *E. coli*, *Lactobacillus spp* and *Salmonella spp*

ranged from 20.12 - 34.98 ( $\log_{10}$ CFU/g), 15.40 - 30.44 ( $\log_{10}$ CFU/g) and 18.20 - 29.09 ( $\log_{10}$ CFU/g). *E. coli* and *Salmonella spp* values were highest in T1 relative to other treatments ( $P < 0.05$ ) contrary to *Lactobacillus spp* count where T5 was highest, T2, T3, T4 followed similar trend and lowest in T1 ( $P < 0.05$ ).

**Table 4.** Caecal microbial population of broiler chicks fed diets with different level of GIGM.

Parameter ( $\log_{10}$ CFU/g)	T1	T2	T3	T4	T5	SEM
<i>E. coli</i>	34.98 <sup>a</sup>	25.10 <sup>b</sup>	23.98 <sup>b</sup>	23.18 <sup>b</sup>	20.12 <sup>b</sup>	1.30
<i>Lactobacillus spp</i>	15.40 <sup>c</sup>	20.76 <sup>b</sup>	23.48 <sup>b</sup>	28.87 <sup>b</sup>	30.44 <sup>a</sup>	1.65
<i>Salmonella</i>	29.09 <sup>a</sup>	19.32 <sup>b</sup>	19.04 <sup>b</sup>	18.58 <sup>b</sup>	18.20 <sup>b</sup>	0.96

Means in the same row with different superscripts differ significantly ( $P < 0.05$ ).

**DISCUSSION**

The pharmacological importance of EOs is primarily due to bioactive chemicals in plant tissues as primary and secondary metabolites [17]. These constituents have several therapeutic properties for instance terpenoids possess anticarcinogenic, antimalarial, anti-ulcer, antimicrobial or diuretic activity [18,19]. Flavonoids in plants possess medicinal benefits which includes antioxidant and anti-inflammatory activities [20,21]. They have the ability to scavenge hydroxyl radicals, super oxide anions and lipid peroxy radicals [22]. Alkaloids perform antimalarial, antimicrobial, antioxidant and protection of plants from pathogens [23]. Phenolic compounds show a wide range of pharmacological activities including anticancer, anti-inflammatory and prevention of cardiovascular diseases [24].

The superior growth performance observed among birds in T3, T4 and T5 ( $P < 0.05$ ) could be attributed to the essentials oils combination which exerted synergistic effects to prevent the consequence of intestinal inflammation. Activities of phytochemicals in GIGM have also proven to stimulate functions of the intestinal tract to improve digestive secretions, nutrient absorption and metabolism [25]. The dietary inclusion of GIGM in broilers also exerted a significant difference in feed intake ( $P < 0.05$ ).

The activities of pathogenic bacteria in the caecum of the birds decreased as the dietary inclusion of GIGM increases ( $P < 0.05$ ) across the treatments. This result agrees with the findings of Singh [1] and Adewale [2] who reported that phytochemicals such as flavonoids, phenols and alkaloids are capable of reducing the activities of pathogenic bacteria through competitive exclusion and promoting the proliferation of beneficial bacteria like *Lactobacillus spp*,

thus playing a role of a probiotic. GIGM can act at different sites of the gastrointestinal tract (GIT) and relay on different targets such as modifying the intestinal microbial balance in favor of beneficial bacterial strains, colonization of the mucosa by adhering non-pathogens and occupation of specific receptors on mucosal surface (prebiotics) [26,27].

**CONCLUSION**

It was concluded from this experiment that the use of essential oils (GIGM) is effective and effective and it represents one of the promising alternatives to antibiotics because it contains several secondary metabolites which performs several medicinal properties. Dietary inclusion of GIGM at 0.4 % had a significant impact on growth as well as reducing the population of pathogenic bacteria without causing any deleterious effect on the performance and health of the animal.

**REFERENCES**

1. Singh AS, Alagbe JO, Sharma S, Oluwafemi RA, Agubosi OCP (2021) Effect of dietary supplementation of melon (*Citrullus linatus*) seed oil on the growth performance and antioxidant status of growing rabbits. J Multidimension Res Rev 2(1): 78-95.
2. Adewale AO, Alagbe JO, Adekemi AO (2021) Dietary Supplementation of *Rauvolfia Vomitoria* Root Extract as A Phytogetic Feed Additive in Growing Rabbit Diets: Hematology and serum biochemical indices. Int J Orange Technol 3(3): 1-12.
3. Musa B, Alagbe JO, Betty AM, Omokore EA (2020) Growth performance, caeca microbial population and immune response of broiler chicks fed aqueous extract

- of *Balanites aegyptiaca* and *Alchornea cordifolia* stem bark mixture. United J Res Technol 2(2): 13-21.
4. Michiels J, Missotten J, Dierick N, De Smet S (2005) *In vitro* effect of botanicals on gut flora of pigs. 30ste Studiedag van de Nederlandstalige Voedingsonderzoekers Merelbeke. pp: 67-68.
  5. Charles R, Garg SN, Kumar S (2000) New gingerdione from the rhizomes of *Zingiber officinale*. Fitoterapia 71: 716-718.
  6. Chang KJ, Cheong SH (2008) Volatile organosulphur and nutrient compounds from garlic by cultivating areas and processing methods. Fed Am Soc Exp Bio J 22: 1108-1112.
  7. Alagbe JO, Oluwafemi RA (2019) Performance and hematological parameters of broiler chicks gives different levels of dried lemon grass (*Cymbopogon citratus*) and garlic (*Allium sativum*) extract. Res Agric Vet Sci 3(2): 102-111.
  8. Cheng S, Liu JY, Cheng EH, Cheng ST (2008) Antifungal activity of cinnamaldehyde and eugenol congeners against wood rots fungi. Bioresour Technol 99: 5145-5149.
  9. Demir E, Sarica S, Ozcan MA, Suicmez M (2003) The use of natural feed additives as alternatives for an antibiotic growth Promoter in broiler diets. Braz J Poult Sci 44: 44-45.
  10. Olafadehan OA, Oluwafemi RA, Alagbe JO (2020) Carcass quality, nutrient retention and caeca microbial population of broiler chicks administered Rolfe (*Daniellia oliveri*) leaf extract as an antibiotic alternative. J Drug Discov 14(33): 146-154.
  11. Dormans HJ, Deans SG (2000) Antimicrobial agents from plants: Antibacterial activity of plant volatile oils. J Appl Microbiol 88: 308-316.
  12. Burt S (2004) Essential oils: Their antibacterial properties and potential applications in food- A review. Int J Food Microbiol 94: 943-947.
  13. Oluwafemi RA, Uankhoba IP, Alagbe JO (2021) Effects of turmeric oil as a dietary supplement on the growth performance and carcass characteristics of broiler chicken. Int J Orange Technol 3(4): 1-9.
  14. Balogun O (2001) The Federal Capital Territory of Nigeria: Geography of Its Development. University of Ibadan Press Limited.
  15. Harborne JD (1973) Phytochemical methods: A guide to modern techniques of plant analysis. Chapman and Hall, London. pp: 279.
  16. Odebiyi A, Sofowora AE (1978) Phytochemical Screening of Nigerian Medicinal Plant. Part III, Lloydia, 41: 234-246.
  17. Shittu MD, Alagbe JO (2020) Phyto-nutritional profiles of broom weed (*Sida acuta*) leaf extract. Int J Integr Educ 3(11): 119-124.
  18. Dudareva N, Pichersky E, Gershenzon J (2004) Biochemistry of plant volatiles. Plant Physiol 135: 1893-1902.
  19. Krishnaiah D, Sarbatly R, Bono A (2007) Phytochemical antioxidants for health and medicine: A move towards nature. J Biomed Sci 1(3): 97-104.
  20. Saxena M, Saxena J, Nema R, Singh D, Gupta A (2013) Phytochemistry of Medicinal Plants. J Pharmacogn Phytochem 8192(1): 168-182.
  21. Ojewuyi OB, Ajiboye TO, Adebajo EO, Balogun A, Mohammed AO (2014) Proximate composition, phytochemical and mineral contents of young and mature *Polyalthia longifolia* Sonn. leaves. Fountain J Nat Appl Sci 3(1): 10-19.
  22. Okwu DE (2004) Phytochemical and vitamin content of indigenous spices of South Eastern Nigeria. J Sustain Agric Environ 6: 30-37.
  23. Cushnie TPT, Cushnie B, Lamb AJ (2014) Alkaloids: An overview of their antibacterial antibiotic enhancing and anti-virulence activities. Int J Phytochem 44(5): 377-386.
  24. Li S, Li RY, Gan FL, Li HB (2013) Antioxidant capacities and total phenolic contents of infusions from 233 medicinal plants. Ind Crop Prod 51: 289-298.
  25. Alagbe J, Oluwafemi RA (2019) Growth performance of weaner rabbits fed noni (*Morinda citrifolia*) and *Moringa olifera* leaf mixture as partial replacement for soya bean meal. Int J Adv Biol Biomed Res 7(2): 185-195.
  26. Michiels J, Missotten J, Dierick N, De Smet S (2005) *In vivo* degradation and *in vivo* passage kinetics of carvacol, thymol, eugenol and trans-cinnamaldehyde along the gastrointestinal tracts of piglets. J Food Agric 88: 2371-2381.
  27. Dierick N, Michiels J, Van NC (2004) Effect of medium chain fatty acids and benzoic acid as alternative for antibiotics on growth and some gut parameters in piglets. Commun Agric Appl Biol Sci 69(2): 187-190.