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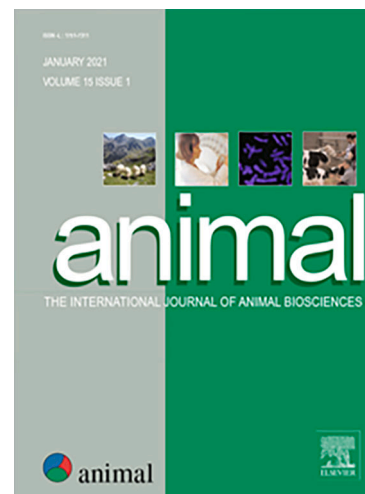
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Sensitivity of broiler performance, organ weights and plasma constituents to amino acid supplementation and reused litter exposure using ideal protein formulated rations

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Abstract

Effects of amino acid supplementation to ideal protein (**IP**) formulated rations were investigated on growth performance, plasma metabolites and organ weights of broilers placed on 100% recycled (reused) litter. Day-old Ross308 male broilers were raised on either clean or reused litter and fed for three weeks on one of five isoenergetic diets, where an IP-based control diet (**C**) was compared with diets containing threonine (**T**) or arginine (**A**) at 25% above requirements, or with 1% supplemented glutamine (**G**), or with each amino acid added (**TAG**). Litter and diet treatments did not strongly interact on outcomes. Reused litter placement resulted in greater weight gain, smaller feed conversion ratio and heavier bursal weights ($P < 0.05$) compared to clean litter placement. Relative to C and T birds, TAG birds reduced weight gain and feed intake ($P < 0.05$). Plasma uric acid levels in G birds were greater than in C, T and A birds ($P < 0.001$). Collectively, since the outcomes of placement on reused litter increased performance and the control diet was IP formulated, the absence of increased growth performance in response to amino acid supplementation would be consistent with amino acids tested being excess to requirements.

Keywords

Threonine, arginine, glutamine, reused poultry litter, growth performance.

Implications

This manuscript explores the sensitivity of using the ideal protein concept to reused litter exposure as a model to mimic litter-borne microbiome exposure in the poultry industry. Our findings confirm that the ideal protein concept remains an appropriate basis for an effective feeding strategy, with no extra benefit from selected amino acid supplementation as reused litter exposure increased performance. Thus, our study supports the view that multiple benefits can arise from reused litter usage, including increased feed and non-feed resource use efficiency, both contributing to reducing environmental footprint, which will be of global interest including Europe.

Introduction

Many studies have shown that supplementing broiler rations with selected essential amino acids (**AA**) enhance performance, including for threonine (Min et al., 2017; Ji et al., 2019; Ahmed et al., 2020) and arginine (Ebrahimi et al., 2014; Pirsaraei et al., 2018; Omid et al., 2020), but also for the non-essential AA glutamine (Nassiri Moghaddam and Alizadeh-Ghamsari, 2013; Ribeiro et al., 2015; Wu et al., 2021), suggesting that availability of these AA for growth performance was reduced from the control rations used. To minimise excess protein supply and excessive N excretion, the ideal protein (**IP**) concept has been introduced in diet formulations (Baker, 2003; Miles and Chapman, 2007; McGill et al., 2012). A diet is formulated to IP if it provides digestible AA in the ratios healthy birds require for maintenance and production (Deschepper and de Groote, 1995). Consequently, increased performance would not be expected from AA supplementation to an IP formulated baseline. However, selected AA supplementation resulted in increased resilience and immune responses when birds were under a wide range of sub-clinical enteric health challenges, including sub-clinical coccidiosis, necrotic enteritis, and exposure to reused litter (Star et al., 2012; Keerqin et al., 2017; Bortoluzzi et al., 2020). These studies suggest that sub-clinical enteric microbial challenge results in different dietary AA ratios for optimal production, and thus may influence the IP concept.

Specific AA are required for effective immune response, including lymphocyte proliferation and the synthesis of immunoglobulins, nitric oxide, and cytokines (Rubin et al., 2007). Among the essential AA, threonine is required to synthesise mucin, which covers the intestinal mucosal surface (Ahmed et al., 2020), and threonine supplementation has resulted in increased antibody responses (Sigolo et al., 2017). During enteric disease challenges, reduced threonine availability may restrict mucin synthesis, impairing gut barrier function. It has been reported that threonine supplementation may enhance mucin synthesis and enhance mucosal integrity during enteric challenges (Faure et al., 2007; Mehdipour et al., 2020). Many studies have demonstrated that threonine supplementation can increase performance and gut health in broilers under enteric challenge (Kidd et al., 2003; Star et al., 2012; Mehdipour et al., 2020). However, others show no such effects (Chen et al., 2017), even in the presence of sub-clinical challenges (Corzo et al., 2007; Valizade et al., 2014).

Supplementing diets with arginine has increased broiler growth performance under commercial conditions, where sub-clinical challenge is assumed to be present to some

degree (Montanhini Neto et al., 2013). In addition, supplementing diets with arginine may reduce intestinal permeability, indicating its protective role when intestinal inflammation and permeability are elevated (Barekatin et al., 2019 and 2021). Moreover, arginine is essential for enterocyte growth and development (Rhoads et al., 2008) and is required to produce nitric oxide, a key component of innate immunity (Zhang et al., 2018). Whilst glutamine is known as non-essential AA, it has been considered a conditionally essential AA at times of stress and enteric challenges (Curi et al., 2007; Wang et al., 2009). Glutamine is the main energy source for immune and intestinal epithelial cells. It is also a precursor of N-acetyl-glucosamine and N-acetyl-galactosamine, major components of mucin (Coster et al., 2004). In addition, glutamine may enhance performance during sub-clinical challenges by acting on inflammatory processes and increasing the intestinal surface area and, thus, nutrient absorption (Xue et al., 2018; Oxford et al., 2019).

Variability in response to AA supplementation likely has a multifactorial basis, including the level of sub-clinical challenge and deviation of baseline AA profile from IP. Here, we assessed the effects of individual or combined supplementation with threonine, arginine and glutamine and reused litter exposure on performance, blood biochemistry and organ weights in broilers where basal rations were formulated to IP. In a parallel paper, we have recently reported concurring responses in caecal microbiome parameters to link to the performance outcome for the combined AA supplementation and reused litter exposure (Hussein et al., 2023). Raising birds on reused litter may have variable outcomes on performance, as it has been associated with decreased (Khattak et al., 2019; González-Ortiz et al., 2021), similar (Yamak et al., 2016; Vieira and Moran, 1999) or increased (Garcés Gudiño et al., 2018) performance relative to birds raised on clean litter. Therefore, we hypothesised that the outcome of AA supplementation to IP formulated basal rations is sensitive to the outcome of reused litter exposure, and that AA supplementation increases performance only when reused litter exposure reduces bird performance.

Material and methods

Housing and experimental design

A total of 800 one-day-old Ross 308 broiler male chickens were used in a 21-day experiment. Upon arrival, chicks were weighed and distributed to 80 experimental pens in a single animal room with 10 birds per pen and 8 pens per treatment. Stocking density, temperature and lighting schedule were maintained as described in Hussein et al. (2023). Birds were *ad libitum* fed and watered. Five dietary treatments and two litter treatments were combined in a 5 × 2 factorial arrangement. The resulting ten treatment combinations were assigned within a complete randomised block design to one of ten pens in eight blocks to account for possible pen location effects within the single animal room used.

Diet treatments

The first diet treatment was the control (**C**), which was IP formulated and included crystalline lysine, methionine, threonine, tryptophan, arginine, valine, and isoleucine to meet all essential AA requirements on a digestible AA basis. The 2nd and the 3rd diet treatment consisted of additional threonine (**T**) and arginine (**A**), respectively, at 25% above requirements. The 4th diet treatment consisted of 1% glutamine

supplementation (**G**). The 5th diet treatment combined additional threonine, arginine and glutamine at the aforementioned levels (**TAG**). The inclusion levels of T, A and G were informed by Corzo et al. (2007), Fasina et al. (2010) and Gottardo et al. (2016), respectively. The tested AA were included in diet C at the expense of maize starch. Diets were calculated to be isoenergetic for apparent metabolisable energy and varying CP levels, as presented in Tables 1 and 2. The experimental diets were fed as mash during starter (day 0 to 11) and grower (day 11 to 21) phases, and were formulated to Ross 308 recommendations (Aviagen, 2014).

Litter treatments

The litter within each pen was either unused wood shavings (clean litter) or with spent litter from a previous broiler study with no evidence of clinical disease (reused litter). The reused litter was stored in bags at room temperature for four weeks between collection and usage.

Chemical analysis of the diets

Representative samples of the experimental diets were analysed for DM, gross energy, CP, and all AA, as described elsewhere (Hussein et al., 2023). The analysed nutrient contents of the experimental diets were mostly consistent with the expected values (Table 3 and Table 4), though analysed CP contents of T, A and G diets were, on average, ~1% smaller than calculated. However, the analysed individual AA contents generally accorded with the calculated figures.

Growth performance

Growth performance parameters, i.e., body weight gain (**BWG**), feed intake (**FI**), and feed conversion ratio (**FCR**, calculated as FI over BWG), were calculated from mean body weights through bulk weighing and bird counting at pen level, and weights of feed offered on days 0 and 11, and feed refusals on days 11 and 21. The resulting BWG, FI and FCR were calculated for the periods of days 0 to 11 (starter phase), 11 to 21 (grower phase), and 0 to 21 (total trial). Dead or culled birds were weighed, and date recorded, to correct performance parameters for mortality.

Blood parameters and organ weights

Two birds per pen were selected randomly at trial end (day 21) in the morning, individually weighed, electrically stunned, and exsanguinated in the post-mortem room (away from home cages) to collect 5 mL of blood in K-ethylenediaminetetraacetic acid treated vacutainers. Plasma was separated through centrifugation at 2000 × g for 15 min at room temperature and was stored at -20 °C pending analysis. To underpin influences of reused litter exposure on performance and its possible modification through AA supplementation, plasma was analyzed for total protein, albumin, uric acid, triglycerides, total cholesterol, total bile acids, glucose, gamma-glutamyl transpeptidase, glutamate dehydrogenase, Na, K and Cl through an AU 5800 clinical chemistries analyzer (Beckman) at Synlab Vet GmbH (Geesthacht, Germany). Plasma globulin was calculated as the difference between the total protein and albumin. In addition, the gizzard with the intact yellow lining membrane, spleen and bursa were removed and weighed. The absolute organ weights were reported to give

an indication of gut development (gizzard) and as associated with the immune system (spleen and bursa).

Litter analysis

Litter samples for both clean and reused litter treatments were collected at day 0 using sterilised gloves and plastic bags. The collected litter samples were mixed thoroughly in sterilised trays, packed inside sealed plastic bags, and kept at $-80\text{ }^{\circ}\text{C}$ pending pH, moisture content and microbiological analysis.

Litter pH and moisture. Litter samples were defrosted, spread onto a clean sheet of plastic, mixed thoroughly and sub-sampled in duplicate for pH and moisture content analysis, as described elsewhere (Hussein et al., 2023). Litter moisture was then calculated from the sample start and end weight difference. The resulting moisture content of the clean and reused litter at day 0 (\pm SE) was $10.8 \pm 2.3\%$ and $12.5 \pm 2.3\%$, respectively, and had a pH of 5.8 ± 0.1 and 8.1 ± 0.1 , respectively.

Microbiological analysis for bacterial detection in collected litter samples. Litter samples at day 0 were tested in duplicate for the presence of *Escherichia coli* and *Campylobacter jejuni* using the culturing method. Strains of *E. coli* and *C. jejuni* used as positive controls were retrieved from frozen beads and stored at $-80\text{ }^{\circ}\text{C}$ at SAC Veterinary Services, Auchincruive, UK.

10 g of litter samples were added to 50 mL sterile phosphate buffer saline in a sterilised stomacher bag and homogenised using a mechanical shaker for approx. 1 min. For *E. coli*, 10 μL of litter suspension were spot dropped onto MacConkey agar plates. The plates were aerobically incubated at $37\text{ }^{\circ}\text{C}$ for 24 h. Isolated colonies were identified as *E. coli* by observing their cultural characterisation, gram's stain, oxidase test and biochemical reaction using indole test (Munoz et al., 2021). Whereas, for *C. jejuni*, 100 μL of the litter mixture was applied onto charcoal cefoperazone deoxycholate agar plates (Thermo-Fisher Scientific) and spread evenly over the surface with a sterile spreader. Plates were incubated invertedly at $41.5\text{ }^{\circ}\text{C} \pm 2.0\text{ }^{\circ}\text{C}$ in an atmosphere suitable for microaerophilic and capnophilic micro-organisms and examined after 48 hours of incubation. For confirmation, two colonies were picked up from each plate and sub-cultured onto blood agar plates (Oxoid PB0114) and incubated at $37\text{ }^{\circ}\text{C}$ for 48 h, one plate aerobically and one plate microaerophilically. The presence of *C. jejuni* was indicated by a lack of growth aerobically and colonies with *C. jejuni* morphology that grew microaerophilically. Gram staining and oxidase strips (Oxoid Ltd., MB0266) were performed on all samples (Khattak et al., 2018). Both clean and reused litter were positive for oxidase-negative and indole-positive *E. coli* but negative for *C. jejuni*.

Statistical analysis

Data collected were analysed using the general ANOVA function of GenStat (16th Edition) and following the 5×2 factorial arrangement for diet treatments (C, T, A, G and TAG), litter treatments (clean vs reused) and their interaction, using pen location as a block and day 0 body weight as a covariate for day 11 and day 21 body weight. Prior to analysis, the data were checked for normality by examining residuals, histograms, and box plots. Data that were not normally distributed were log₁₀ transformed prior to statistical analysis. Means presented without common superscript differed significantly as per Tukey's honest significance test at $P < 0.05$. Power analysis

supported that at a 2% coefficient of variation for BWG and FCR, typical for poultry trials at SRUC, Auchincruive, biologically meaningful differences of 3.5% in response to treatment were expected to be detected at $n = 8$.

Results

Growth performance and organ weights

Tables 5a and 5b show the effect of litter and diet treatments on FI. During the starter phase, litter and diet treatments did not interact. Birds placed on reused litter ate less than birds placed on clean litter, whilst TAG birds ate less than C and T birds, with A and G birds being intermediate. During grower phase and total trial, litter and diet treatments interacted; TAG birds ate less than C and A birds placed on clean litter, whilst diet did not affect FI of birds placed on reused litter.

Table 6 shows the effect of litter and diet treatments on BWG and FCR. Initial body weight (day 0) did not differ significantly between treatments ($P=0.265$) and averaged (\pm SD) 39.3 ± 0.81 g. Litter and diet treatments did not interact on BWG and FCR. However, birds placed on reused litter had greater BWG and smaller FCR than birds placed on clean litter, whilst TAG birds had smaller BWG and greater FCR than C and T birds, with A and G birds being intermediate and the effect on FCR limited to the starter phase. Mortality was low at 0.5% (4 out of 800 birds placed) and was not treatment associated.

Table 6 also shows the effect of litter and diet treatments on empty gizzard, spleen, and bursa weights. There were no main effects of diet treatment or interaction between diet and litter treatments on organ weights. However, litter treatments independently affected organ weights, as birds placed on reused litter had a lighter empty gizzard and a heavier bursa than birds placed on clean litter, with no effect on spleen weight.

Plasma parameters

Most read-outs on plasma biochemistry were not affected by the experimental treatments, which averaged (\pm SD) 29.5 ± 2.75 g/L for total protein, 12.2 ± 1.20 g/L for albumin, 17.3 ± 1.73 g/L for globulin, 19.0 ± 3.70 U/L for gamma-glutamyl transpeptidase, 18.9 ± 7.58 μ mol/L for bile acids, 3.6 ± 0.52 mmol/L for cholesterol, 1.6 ± 0.46 mmol/L for triglycerides, 16.1 ± 1.03 mmol/L for glucose and 95 ± 7.9 mmol/L for Cl. However, diet treatments influenced plasma uric acid, which averaged 444^{ab} , 436^{ab} , 369^a , 553^c and 520^{bc} μ mol/L for C, T, A, G and TAG birds, respectively (SED 38.7 mmol/L; $P<0.001$). In addition, litter treatments influenced plasma Na and K concentrations. Plasma Na averaged 143.7 and 138.2 mmol/L for birds placed on clean and reused litter, respectively (SED 2.63 mmol/L; $P<0.05$), whilst their plasma K averaged 27.1 and 24.5 mmol/L, respectively (SED 0.86 mmol/L; $P<0.05$). Gamma-glutamyl transpeptidase averaged 4.3 ± 1.23 U/L, and although diet and litter treatment interacted ($P=0.046$), Tukey's honest significance test at $P<0.05$ revealed no diet treatment effect within litter treatment, or vice versa.

Discussion

Here, we assessed the effect of specific AA supplementation to IP formulated rations on growth performance, blood biochemistry, and organ weights of birds placed on

clean or reused litter. Since the control diet was formulated to IP, it was hypothesised that the effect of AA supplementation is sensitive to the outcomes of reused litter exposure. Our data support the hypothesis, as having observed that reused litter increased bird performance, AA supplementation reduced the growth performance of birds in both litter treatments.

The results support the view that birds fed C diets had greater BWG, FI and smaller FCR than those fed TAG diets. The C diets were formulated on digestible AA requirements (IP-basis) to maximise growth performance and reduce N excretion into the environment (Adedokun et al., 2016; Maharjan et al., 2020). Our results showed that the individual T, A, and G diets tended to reduce bird performance. It can be proposed that these relatively small individual negative effects accumulated in the larger penalty on performance when these AA were combined in the TAG diet. The latter could be attributed to multiple causes. Firstly, diets were formulated to be isoenergetic, whilst threonine, arginine, and glutamine supplementation in TAG diets increased the CP content. This greater CP level above IP needs to be metabolised, requiring additional energy use for the process of excess AA deamination and N elimination. It has been estimated that 6 molecules of ATP are used to excrete each excess N molecule, which contrasts with the 4 molecules of ATP required per N molecule for protein synthesis (Aletor et al., 2000). This accords with observations that a reduced apparent metabolisable energy to CP ratio reduces energy available for growth to benefit uric acid synthesis (Dessimoni et al., 2019) and agrees with elevated levels of uric acid observed here. The latter may also indicate an enhanced caecal protein and/or AA fermentation (Elling-Staats et al., 2022), a view that is supported by elevated levels of caecal branched chain fatty acids in TAG birds (Hussein et al., 2023). Secondly, an increased N excretion from excess AA supplementation might result in elevated ammonia production in the litter, e.g. through uric acid degradation, which may detriment broiler health, environment and performance (Aletor et al., 2000; Corzo et al., 2005). It should also be recognised that the final body weight was 20% below Ross 308 target (Aviagen, 2014). This is likely due to mash feeding rather than crumbs (starter) and pellets (grower). Compared to mash, pelleted feeds usually result in enhanced bird performance arising from amongst others decreased feed wastage, increased feed intake, avoidance of selective feeding, pathogen reduction, improved palatability, and increased nutrient digestibility (Lv et al., 2015).

Raising birds on reused litter has shown variable outcomes on performance, as it has been associated with decreased (Torok et al., 2009; Khattak et al., 2019; González-Ortiz et al., 2021), similar (Yamak et al., 2016; Vieira and Moran, 1999) or increased (Garcés Gudiño et al., 2018) performance relative to birds raised on clean litter. It might be speculated that such variability in response to reused litter exposure could result from variations in pathogen load, moisture, pH, and recycled nutrients from the previous flock. In our study, birds on reused litter had 4.5% greater final body weight than those on clean litter by day 21. Gut-derived microbiome in the reused litter may have acted as a probiotic or a direct-fed microbial supplementation that can alter the gastrointestinal tract microbiota of newly hatched chicks, leading to enhanced performance (Torok et al., 2009; Shanmugasundaram et al., 2012). In addition, the more alkaline pH of reused litter used here may indicate greater ammonia production, which may influence pathogen load, as indicated from earlier studies (Kennard et al., 1959). Furthermore, reused litter microbiome has been shown to reduce caecal abundance of *Salmonella* spp. and delay *Clostridium perfringens* colonisation with a

potential increase in broiler performance (Cressman et al., 2010; Wei et al., 2013). This view is supported by a shift towards more beneficial caecal microbiome composition for the birds placed on reused litter (Hussein et al., 2023). Finally, although the rations used were supplemented with vitamin pre-mix, the reused litter might have been a source of vitamins with greater bioavailability, which could also have increased broiler performance. Earlier studies showed that synthesis of vitamin B12 and riboflavin by microorganisms found in chicken droppings increased bird's growth and feed efficiency when raised on reused litter (Kennard et al., 1959; Roll et al., 2011). The reduced FI of birds on reused litter during the starter phase concurred with a smaller FCR, which suggests that ingesting reused litter competes with the feed offered. This supports previous evidence by Malone (1992), who showed that broiler chickens may consume up to 4% of the daily intake as litter. Although there was no significant effect of reused litter on FI in the grower phase or the whole growth period, by day 21, birds on reused litter had a significantly greater body weight, BWG, and smaller FCR compared to birds on clean litter. Since reused litter exposure increased performance, AA supplementation would have resulted in excess AA and as such did not increase bird's performance despite the two contrasting litter treatments used.

Gizzard weight was recorded as a marker for gut development, including in response to variation in intake, whilst spleen (primary lymphoid organ) and bursa (secondary lymphoid organ) weights were recorded to infer possible influence on immune system development (Madej et al., 2015). Diet treatments did not influence gizzard weight, though gizzards of reused litter birds were lighter than those on clean litter, which concurred with reduced FI. Although mash diets were used, which may require less gizzard action than pelleted diets (Svihus, 2011), using the latter would likely have resulted in similar effects on gizzard weight due to its sensitivity to variation in feed intake per se.

The heavier bursa of Fabricius arising from raising birds on reused litter observed may be an indicator of B-lymphocyte development (Rajput et al., 2013). This would be consistent with previous studies where early exposure to the reused litter microbiome increased antigens' subsequent uptake through cloacal reflex, stimulated B-cell development, and resulted in an effective antibody response (Sorvari et al., 1975). On the other hand, there was no significant effect of AA supplementation on bursa weight. Whilst our outcome agrees with Xu et al. (2018), a heavier bursa was reported in birds following supplementation with arginine (Toghyani et al., 2019) and glutamine (Ribeiro et al., 2015), where production responses were also observed. This would further support the view that AA availability in the current study was unlikely to be scarce in the absence of AA supplementation. Spleen weight was not affected in our study, which accords with other studies on threonine supplementation (Toghyani et al., 2019) or arginine and glutamine supplementation (Szabó et al., 2014).

Blood biochemical can assist in evaluating bird health and metabolic status, and how these respond to internal and external factors (Abdul Basit et al., 2020). Most of the measured plasma parameters in the current study were within the reference range of broilers, supporting the observation that placement on reused litter did not detriment the birds and that homeostatic mechanisms were operating within normal physiological range in response to the AA supplementation. Indeed, whilst plasma uric acid levels are also within a normal range (Clinical Diagnostic Division, 1990), they seem to be greater with G supplementation, both in G and TAG birds, which agrees

with the excess nature of the AA supplemented. Only plasma K was elevated beyond normal values, which may have resulted from using K-ethylenediaminetetraacetic acid treated tubes (Cornes et al., 2008).

Conclusion

Placement on reused litter increased body weight gain and reduced feed conversion ratio compared to placement on clean litter. Because of this outcome, and since the control diet was based on the ideal AA profile, the absence of increased growth response to specific AA supplementation would be consistent with AA being excess.

Ethics approval

All the experimental animal procedures in the current study were carried out under the Animals [Scientific Procedures] Act (1986) and approved by SRUC's Animal Welfare and Ethical Review Body (AU AE 33-2018) and carried out under Home Office authorisation (PPL P32D394C9). All methods were carried out in accordance with the Code of Recommendations for the Welfare of Livestock: Meat Chickens and Breeding Chickens (Department for Environment, Food & Rural Affairs, 2018).

Data and model availability statement

The datasets generated during and/or analysed during the current study and the models used are available from the corresponding author upon reasonable request.

Declaration of Generative AI and AI-assisted technologies in the writing process

The authors did not use any artificial intelligence assisted technologies in the writing process

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Author contributions

MAH, FK, LV, SA, JGMH contributed to experimental design; MAH contributed to carrying out the experiment, data analysis, writing-original draft, and presentation of the published work; FK and JGMH contributed to the provision of study materials; FK, LV, SA and JGMH contributed to review & editing and supervision; JGMH contributed to project administration and funding acquisition. All authors read and approved the final version of the manuscript and approved publication.

Declaration of interests

None.

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Table 1 Feed ingredients and as fed calculated chemical compositions of the experimental starter rations fed to male broilers.

	Experimental starter rations				
	C ¹	T	A	G	TAG
Ingredients (%)					
Wheat	58.22	58.22	58.22	58.22	58.22
Maize starch	3.00	2.78	2.66	2.00	1.44
Soybean meal	31.59	31.59	31.59	31.59	31.59
Soya oil	2.20	2.20	2.20	2.20	2.20
Salt	0.05	0.05	0.05	0.05	0.05
Limestone	0.95	0.95	0.95	0.95	0.95
Dicalcium Phosphate	1.85	1.85	1.85	1.85	1.85
Sodium bicarbonate	0.50	0.50	0.50	0.50	0.50
L-Lysine HCl	0.39	0.39	0.39	0.39	0.39
DL-Methionine	0.24	0.24	0.24	0.24	0.24
L-Threonine	0.23	0.45	0.23	0.23	0.45
L-Valine	0.09	0.09	0.09	0.09	0.09
L-Tryptophan	0.14	0.14	0.14	0.14	0.14
L-Arginine	0.10	0.10	0.44	0.10	0.44

L-Isoleucine	0.05	0.05	0.05	0.05	0.05
L-Glutamine	0.00	0.00	0.00	1.00	1.00
Vitamin & Mineral premix ²	0.40	0.40	0.40	0.40	0.40
Calculated chemical composition ³					
Dry matter (%)	88.80	88.80	88.80	88.80	88.80
Crude protein (%)	22.47	22.63	23.16	23.67	24.51
Metabolisable energy (MJ/kg)	12.51	12.51	12.51	12.51	12.51
Ca (%)	0.96	0.96	0.96	0.96	0.96
P (%)	0.72	0.72	0.72	0.72	0.72
Available P %	0.48	0.48	0.48	0.48	0.48
Na (%)	0.19	0.19	0.19	0.19	0.19
Cl (%)	0.16	0.16	0.16	0.16	0.16
Digestible amino acids (%)					
Arginine	1.37	1.37	1.71	1.37	1.71
Histidine	0.48	0.48	0.48	0.48	0.48
Isoleucine	0.86	0.86	0.86	0.86	0.86
Leucine	1.4	1.4	1.4	1.4	1.4
Lysine	1.28	1.28	1.28	1.28	1.28
Methionine	0.51	0.51	0.51	0.51	0.51

Cysteine	0.31	0.31	0.31	0.31	0.31
Tryptophan	0.20	0.20	0.20	0.20	0.20
Threonine	0.86	1.08	1.08	0.86	1.08
Valine	0.96	0.96	0.96	0.96	0.96
Methionine + Cysteine	0.82	0.82	0.82	0.82	0.82
Phenylalanine + Tyrosine	1.58	1.58	1.58	1.58	1.58

¹ C, ideal protein formulated control diet; T, threonine supplemented diet; A, arginine supplemented diet; G, glutamine supplemented diet; TAG, threonine, arginine, and glutamine supplemented diet.

² Vitamin and mineral premix provided (units kg⁻¹ diets): vitamin A (retinyl acetate) 16 000 iu; vitamin D3 (cholecalciferol) 3 000 iu; vitamin E (dl- α -tocopherol acetate) 75 iu; vitamin B1 (thiamin) 3 mg; vitamin B2 (riboflavin) 10 mg; vitamin B6 (pyridoxine HCl) 3 mg; vitamin B12 (cyanocobalamin) 15 μ g; phylloquinone 5 mg; nicotinic acid 60 mg; pantothenic acid 14.5 mg; folic acid 1.5 mg; biotin 275 μ g; choline chloride 250 mg; iron 20 mg; copper 10 mg; manganese 100 mg; cobalt 1 mg; zinc 82 mg; iodine 1 mg; selenium 0.2 mg; molybdenum 0.5 mg.

³ Derived from Premier Atlas data (2014).

Table 2 Feed ingredients and as fed calculated chemical compositions of the experimental grower rations fed to male broilers.

Ingredients (%)	Experimental grower rations				
	C ¹	T	A	G	TAG
Wheat	60.56	60.56	60.56	60.56	60.56
Maize starch	3.00	2.81	2.70	2.00	1.51
Soybean meal	28.50	28.50	28.50	28.50	28.50
Soya oil	3.50	3.50	3.50	3.50	3.50
Salt	0.05	0.05	0.05	0.05	0.05
Limestone	0.87	0.87	0.87	0.87	0.87
Dicalcium Phosphate	1.65	1.65	1.65	1.65	1.65
Sodium bicarbonate	0.50	0.50	0.50	0.50	0.50
L-Lysine HCl	0.32	0.32	0.32	0.32	0.32
DL-Methionine	0.21	0.21	0.21	0.21	0.21
L-Threonine	0.18	0.37	0.18	0.18	0.37
L-Valine	0.06	0.06	0.06	0.06	0.06
L-Tryptophan	0.12	0.12	0.12	0.12	0.12
L-Arginine	0.05	0.05	0.35	0.05	0.35

L-Isoleucine	0.03	0.03	0.03	0.03	0.03
L-Glutamine	0.00	0.00	0.00	1.00	1.00
Vitamin & Mineral premix ²	0.40	0.40	0.40	0.40	0.40
Calculated chemical composition ³					
Dry matter (%)	88.85	88.85	88.85	88.85	88.85
Crude protein (%)	21.01	21.15	21.61	22.21	22.95
Metabolisable energy (MJ/kg)	12.94	12.94	12.94	12.94	12.90
Ca (%)	0.87	0.87	0.87	0.87	0.87
P (%)	0.67	0.67	0.67	0.67	0.67
Available P (%)	0.44	0.44	0.44	0.44	0.44
Na (%)	0.19	0.19	0.19	0.19	0.19
Cl (%)	0.15	0.15	0.15	0.15	0.15
Digestible amino acids (%)					
Arginine	1.23	1.23	1.53	1.23	1.53
Histidine	0.45	0.45	0.45	0.45	0.45
Isoleucine	0.78	0.78	0.78	0.78	0.78
Leucine	1.31	1.31	1.31	1.31	1.31
Lysine	1.15	1.15	1.15	1.15	1.15
Methionine	0.47	0.47	0.47	0.47	0.47

Cysteine	0.30	0.30	0.30	0.30	0.30
Tryptophan	0.19	0.19	0.19	0.19	0.19
Threonine	0.77	0.96	0.77	0.77	0.96
Valine	0.87	0.87	0.87	0.87	0.87
Methionine + Cysteine	0.77	0.77	0.77	0.77	0.77
Phenylalanine + Tyrosine	1.48	1.48	1.48	1.48	1.48

¹ C, ideal protein formulated control diet; T, threonine supplemented diet; A, arginine supplemented diet; G, glutamine supplemented diet; TAG, threonine, arginine, and glutamine supplemented diet.

² Vitamin and mineral premix provided (units kg⁻¹ diets): vitamin A (retinyl acetate) 16 000 iu; vitamin D3 (cholecalciferol) 3 000 iu; vitamin E (dl- α -tocopherol acetate) 75 iu; vitamin B1 (thiamin) 3 mg; vitamin B2 (riboflavin) 10 mg; vitamin B6 (pyridoxine HCl) 3 mg; vitamin B12 (cyanocobalamin) 15 μ g; phylloquinone 5 mg; nicotinic acid 60 mg; pantothenic acid 14.5 mg; folic acid 1.5 mg; biotin 275 μ g; choline chloride 250 mg; iron 20 mg; copper 10 mg; manganese 100 mg; cobalt 1 mg; zinc 82 mg; iodine 1 mg; selenium 0.2 mg; molybdenum 0.5 mg.

³ Derived from Premier Atlas data (2014).

Table 3 Analysed chemical composition, gross energy, and amino acids of the experimental starter diets fed to male broilers.

	Experimental starter rations				
	C ¹	T	A	G	TAG
Chemical composition (as fed)					
DM (%)	88.00	88.00	88.00	88.10	89.00
CP (%)	22.82	22.32	22.06	23.83	24.12
Gross energy (MJ/kg)	16.45	16.39	16.53	16.53	16.63
Amino acids (%)					
Methionine	0.55	0.52	0.50	0.51	0.50
Cysteine	0.36	0.37	0.34	0.37	0.36
Methionine + Cysteine	0.90	0.89	0.84	0.88	0.86
Lysine	1.45	1.43	1.26	1.39	1.31
Threonine	0.98	1.12	0.91	0.97	1.16
Arginine	1.48	1.56	1.64	1.50	1.76
Isoleucine	0.95	0.99	0.85	0.96	0.90
Leucine	1.55	1.63	1.38	1.57	1.49
Valine	1.10	1.13	0.98	1.10	1.02
Histidine	0.52	0.55	0.47	0.53	0.50

Phenylalanine	1.02	1.07	0.90	1.03	0.98
Glycine	0.87	0.93	0.78	0.89	0.85
Serine	1.04	1.09	0.92	1.06	1.02
Proline	1.33	1.38	1.22	1.37	1.33
Alanine	0.88	0.93	0.78	0.90	0.85
Aspartic acid	2.04	2.17	1.76	2.08	1.95
Glutamic acid	4.31	4.49	3.98	5.19	5.19

¹ C, ideal protein formulated control diet; T, threonine supplemented diet; A, arginine supplemented diet; G, glutamine supplemented diet; TAG, threonine, arginine, and glutamine supplemented diet.

Table 3 Analysed chemical composition, gross energy, and amino acids of the experimental starter diets fed to male broilers.

	Experimental grower rations				
	C ¹	T	A	G	TAG
Chemical composition (as fed)					
DM (%)	88.30	88.60	88.40	88.20	88.30
CP (%)	21.14	20.29	20.56	20.51	23.21
Gross energy (MJ/kg)	16.77	16.75	16.68	16.49	16.77
Amino acids (%)					
Methionine	0.46	0.46	0.45	0.41	0.50
Cysteine	0.35	0.33	0.36	0.34	0.35
Methionine + Cysteine	0.81	0.79	0.81	0.74	0.85
Lysine	1.26	1.16	1.25	1.07	1.35
Threonine	0.87	1.03	0.87	0.79	1.10
Arginine	1.36	1.25	1.62	1.20	1.68
Isoleucine	0.88	0.81	0.89	0.78	0.90
Leucine	1.47	1.35	1.49	1.32	1.50
Valine	0.99	0.91	1.02	0.90	1.02
Histidine	0.50	0.46	0.51	0.45	0.51

Phenylalanine	0.97	0.89	0.99	0.87	1.00
Glycine	0.84	0.76	0.85	0.76	0.84
Serine	0.99	0.92	1.00	0.90	1.01
Proline	1.27	1.21	1.32	1.25	1.33
Alanine	0.84	0.76	0.85	0.75	0.86
Aspartic acid	1.91	1.72	1.95	1.65	1.97
Glutamic acid	4.17	3.91	4.21	4.67	5.16

¹ C, ideal protein formulated control diet; T, threonine supplemented diet; A, arginine supplemented diet; G, glutamine supplemented diet; TAG, threonine, arginine, and glutamine supplemented diet.

Table 5a Main effects for average feed intake (g) during starter phase (day 0 to 11), grower phase (day 11 to 21) or total trial (day 0 to 21) of male broilers placed as day-old on either clean or reused litter and fed five different diets^{1,2}

	Litter			Diet ³						P-values		
	Control	Reused	SED	C	T	A	G	TAG	SED	Litter	Diet	Litter × Diet
Feed intake (g)												
Starter phase	231 ^a	225 ^b	2.4	238 ^a	230 ^{ab}	226 ^{bc}	228 ^{ab}	217 ^c	3.9	0.030	<0.001	0.528
Grower phase	752	762	9.7	793	754	763	756	719	15.3	0.286	<0.001	0.035
Total trial	711 ^b	744 ^a	11.1	1 032	985	989	984	936	17.8	0.657	<0.001	0.048

¹ Each litter-diet combination was allocated to 8 pens of 10 birds per pen.

² Means within the same row for each model main effect outcome differ at $P < 0.05$ when presented with different superscripts; see Table 5b for such differences between simple means in the presence of significant litter × diet interactions.

³ C, ideal protein formulated control diet; T, threonine supplemented diet; A, arginine supplemented diet; G, glutamine supplemented diet; TAG, threonine, arginine, and glutamine supplemented diet.

Table 5b Simple means for average feed intake (g) during starter phase (day 0 to 11), grower phase (day 11 to 21) or total trial (day 0 to 21) of male broilers placed as day-old on either clean or reused litter and fed five different diets^{1,2}

	Clean litter					Reused litter					
	C ³	T	A	G	TAG	C	T	A	G	TAG	
Feed intake (g)											
Starter phase ⁴											
Grower phase	802 ^a	745 ^{abc}	781 ^{ab}	731 ^{bc}	702 ^c	785 ^{ab}	764 ^{abc}	746 ^{abc}	781 ^{ab}	736 ^{abc}	21.6
Total trial	1 047 ^a	977 ^{abc}	1 011 ^{ab}	960 ^{bc}	919 ^c	1 017 ^{ab}	993 ^{abc}	967 ^{abc}	1 007 ^{ab}	952 ^{bc}	25.0

¹ Each litter-diet combination was allocated to 8 pens of 10 birds per pen.

² Means within the same row for each model outcome presented with different superscripts differ at P<0.05.

³ C, ideal protein formulated control diet; T, threonine supplemented diet; A, arginine supplemented diet; G, glutamine supplemented diet; TAG, threonine, arginine, and glutamine supplemented diet.

⁴ Simple means during starter phase are not shown as litter × diet interaction was not significant (Table 5a).

Table 6 Average body weight gain and feed conversion ratio during starter phase (day 0 to 11), grower phase (day 11 to 21) and total trial (day 0 to 21), and day 21 organ weight (g) of male broilers placed as day-old on either clean or reused litter and fed five different diets^{1,2}

	Litter			Diet ³						P-values		
	Control	Reused	SED	C	T	A	G	TAG	SED	Litter	Diet	Litter × Diet
Body weight gain (g)												
Starter phase	196 ^b	203 ^a	3.0	212 ^a	205 ^{ab}	198 ^{bc}	197 ^{bc}	186 ^c	4.7	0.028	<0.001	0.628
Grower phase	515 ^b	541 ^a	8.7	542	535	532	524	505	13.8	0.004	0.108	0.914
Total trial	711 ^b	744 ^a	11.1	754 ^a	741 ^a	730 ^{ab}	721 ^{ab}	691 ^b	17.7	0.004	0.013	0.913
Feed conversion ratio												
Starter phase	1.18 ^a	1.11 ^b	0.009	1.13 ^b	1.13 ^b	1.15 ^{ab}	1.16 ^{ab}	1.17 ^b	0.015	<0.001	0.010	0.396
Grower phase	1.47 ^a	1.41 ^b	0.026	1.47	1.41	1.44	1.44	1.44	0.041	0.029	0.798	0.504
Total trial	1.42 ^a	1.36 ^b	0.021	1.40	1.36	1.39	1.39	1.39	0.032	0.004	0.786	0.497

Organ weight (g)

Empty gizzard	20.93 ^a	20.04 ^b	0.404	20.11	20.23	20.52	20.22	21.34	0.639	0.028	0.295	0.317
Spleen	0.774	0.789	0.040	0.825	0.841	0.772	0.697	0.772	0.064	0.710	0.186	0.501
Bursa of Fabricius	1.800 ^b	1.995 ^a	0.068	1.988	1.888	1.975	1.875	1.763	0.107	0.005	0.225	0.502

¹ Each litter-diet combination was allocated to 8 pens of 10 birds per pen.

² Means within the same row for each model outcome presented with different superscripts differ at $P < 0.05$.

³ C, ideal protein formulated control diet; T, threonine supplemented diet; A, arginine supplemented diet; G, glutamine supplemented diet; TAG, threonine, arginine, and glutamine supplemented diet.

Highlights

- Placing broilers on reused litter may mimic litter-borne microbiome exposure

- Growth response to amino acids was tested in such birds fed ideal protein rations
- Reused litter exposure increased weight gain and reduced feed conversion ratio
- Threonine, arginine and glutamine supplementation reduced growth performance
- Reused litter usage may benefit production efficiency of ideal protein fed broilers

Journal Pre-proofs