

Dried rose (*Rosa damascena* Mill.) dreg: an alternative litter material in broiler production

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Abstract

The present study was carried out to determine the effects of using dried rose dreg (DRD) as an alternative litter material for broiler performance and microbiological characteristics of litter. A total of 225 day-old broiler chicks was raised on pine wood shavings (PS), DRD and PS+DRD until 42 days. The effects of litter material on broiler performance and FCR did not differ. At 42 days the effects of DRD on microbiological characteristics differed statistically from the other treatments. Total aerobic mesophilic bacteria (TAMB), Enterococci, *Enterobacteriaceae* and *Staphylococcus aureus* counts were decreased in the DRD group at 42 days. These results indicate that DRD can be used as an alternative litter material to control microbial characteristics without a negative effect on broiler performance.

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Introduction

There are numerous challenges presented when attempting to reduce human exposure to foodborne pathogens, especially when these pathogens are associated with foods of animal origin. Reduction of pathogens on farms is necessary if cleaner products are expected after processing. However, few effective intervention methods are currently available for use in poultry production. Several poultry litter treatments such as aluminium sulphate (alum) and sodium bisulphate, intended primarily for use in reducing ammonia levels, promoting animal health or inhibiting mould growth, do exist. Since many of these products are acidic compounds it is reasonable to expect that proper application could significantly lower pH and water activity of the poultry litter, conditions that directly affect the survival rate of microorganisms present in the litter (Line, 2002).

It is well known that ammonia build-up adversely affects body weight, feed conversion and survival rate in poultry and is also harmful to farm labourers. Acidic poultry litter treatments reduce volatile ammonia by reducing the pH, which shifts the $\text{NH}_4^+/\text{NH}_3$ equilibrium towards NH_4^+ (Charles & Payne, 1966; Donham, 1996; Line, 2002). Significant reductions in litter pH and exposure to pH levels below 4.5 would be expected to be bacteriostatic or bactericidal for *Salmonella* and *Campylobacter* (Hill *et al.*, 1995; Line, 2002).

Rose (*Rosa damascena* Mill.) is a species being used to produce attar of rose by distilling volatile oils from the flowers. Turkey is amongst the leading producers of rose oil. The rose oil is produced mainly in Isparta, Burdur and Denizli provinces. One unit weight of rose oil distilled gives about 2 units of residue on a wet basis. The content of solids in the wet residue is approximately 10%, consisting of 90% organic matter (Tosun *et al.*, 2002). It was reported that some fresh and spent flower extracts and essential oils obtained from plants such as *Rosa damascena*, *Mentha piperita*, *Origanum onites*, *Juniperus exalsa*, *Thymbra spicata*, *Satureja cuneifolia* have antioxidant, antimicrobial and antibacterial activities (Aridogan *et al.*, 2002; Baydar *et al.*, 2004; Ozkan *et al.*, 2004).

The present study was conducted to determine the effects of using dried rose dreg alone or equally mixed with pine wood shavings as an alternative litter material in broiler production on broiler performance and microbiological composition of the litter during the production period.

Materials and Methods

A total of 225 day-old Cobb 500 broiler chicks was used for this study conducted at the S. Demirel University, Atabey (Isparta) in Turkey. Chicks were randomly distributed into three trial groups with three replicates of each group. In the control group, broilers were raised on pine wood shavings (PS). In the second and third groups, broilers were raised on DRD and equally mixed PS and DRD litter material (PS+DRD),

respectively. All the broilers were housed in nine enclosed pens with a stocking density of approximately 14 birds/m² for 42 days. The broilers received food and water *ad libitum*.

Litter pH levels were monitored weekly by placing about 25 g samples (three replicates for each group) in 125 mL deionised water and measuring the pH with a pH meter (WTW PH526, Germany). Litter samples were aseptically collected to determine microbiological characteristics at 0, 21 and 42 days. Then, 10 g of litter samples were homogenized with 90 mL of a sterile solution of 0.85% (w/v) sodium chloride at 45 °C. Decimal dilutions (10⁻¹, 10⁻², 10⁻³ or 10⁻⁴) of litter samples were prepared in 9 mL sterile NaCl (0.85%) to count pathogenic microorganisms.

Total aerobic mesophilic bacteria (TAMB) were enumerated on plate count agar following the pour-plate method and incubated at 30 °C for 72 h. *Salmonella* and *Shigella* were grown on S.S. agar incubated at 37 °C for 48 h. *Enterobacteriaceae* and coliforms were determined on Violet Red Bile Glucose agar (VRBG) and Violet Red Bile Lactose (VRBL), respectively and incubated at 37 °C for 24-48 h. *Staphylococcus aureus* was enumerated by plating on Baird Parker egg yolk tellurite medium following the surface-plate method, and incubated at 37 °C for 48 h. Black colonies with cream-white halo and with a clear zone around it were recorded as *S. aureus*. Yeast and moulds were determined on potato dextrose agar acidified with 10% lactic acid following the surface plate method, and incubated at 25 °C for four days. Enterococci were counted on KF Streptococcal agar incubated at 30 °C for three days (APHA, 1995; Sparks, 1996). The microbial counts were determined as colony forming units (cfu) per gram of the samples. The results of microbiological analyses are shown as log cfu/g. All analyses were conducted in duplicate. Statistical analyses of the data were performed using an analysis of variance with separation of the means by the Duncan Multiple Range test (Duzgunes *et al.*, 1987). Statements of statistical differences were based on P ≤ 0.05, unless otherwise stated. All analyses were performed using SPSS[®] 10.0 computer software (SPSS Inc., Chicago, USA 1999).

Results and Discussion

Live weights and feed conversion ratio (FCR) values are shown in Table 1 and Table 2, respectively.

Table 1 Mean weekly body weights, (g ± s.e.) of the chickens raised on different litter material

Weeks	Litter material		
	PS	DRD	PS+DRD
0	41.1 ± 0.62	42.1 ± 0.69	42.1 ± 0.55
1	120.1 ± 2.36	126.6 ± 2.35	123.4 ± 2.11
2	349.7 ± 7.10	362.9 ± 5.78	353.9 ± 5.51
3	753.0 ± 13.7	767.4 ± 11.1	760.4 ± 10.9
4	1287.0 ± 23.0	1321.6 ± 19.9	1299.6 ± 19.8
5	1874.7 ± 34.6	1923.6 ± 31.8	1900.8 ± 23.6
6	2338.8 ± 40.4	2394.2 ± 31.0	2371.1 ± 31.3

PS - pine wood shavings; DRD - dried rose dreg

Table 2 Cumulative feed conversion ratios (FCR) (g feed/g weight gain) (mean ± s.e.) of chickens raised on different litter material

Weeks	Litter material		
	PS	DRD	PS+DRD
0-1	1.003 ± 0.072	0.972 ± 0.033	0.966 ± 0.033
0-2	1.269 ± 0.049	1.207 ± 0.031	1.215 ± 0.015
0-3	1.393 ± 0.046	1.365 ± 0.008	1.338 ± 0.007
0-4	1.549 ± 0.029	1.534 ± 0.019	1.515 ± 0.005
0-5	1.751 ± 0.040	1.675 ± 0.011	1.699 ± 0.009
0-6	1.883 ± 0.031	1.816 ± 0.010	1.838 ± 0.008

PS - pine wood shavings; DRD - dried rose dreg

Results from Table 1 show that up to 42 days of age the body weights of the chickens did not differ ($P > 0.05$) between groups. Nevertheless, the broilers raised on DRD were heavier than other litter treatments throughout the trial. Similarly, birds raised on PSD+DRD were found numerically heavier than the birds raised on PS. According to the cumulative FCR comparison in Table 2, there was no statistical difference ($P > 0.05$) between groups throughout the trial, but the results may be interpreted that FCR was numerically better in DRD litter group than in the other groups.

The microbiological characteristics of litter materials are summarized in Table 3. While initial TAMB and Enterococci counts were lower ($P < 0.001$) in the PS group than in the DRD and PS+DRD groups, it was found that the lowest ($P < 0.05$) TAMB and Enterococci counts were in the DRD group at 42 days. Similarly, the *Enterobacteriaceae* count did not differ until day 21, when there was a lower ($P < 0.05$) *Enterobacteriaceae* count in the DRD group at 42 d than in the other groups. The *S. aureus* count differed ($P < 0.05$) between groups only at 42 days. While initial total yeast and mould counts were found lower ($P < 0.001$) in PS group, it was found at a numerically lower count in the DRD group at 42 days than in the other groups. As an alternative litter material, DRD had antimicrobial and antibacterial effects at 42 days, presumably due to the essential oils. This is in agreement with other researchers (Aridogan *et al.*, 2002; Baydar *et al.*, 2004; Ozkan *et al.*, 2004).

Table 3 Microbiological characteristics in the different litter material, log cfu/g (mean \pm s.e.)

Microbial result	Litter Material	Days		
		0*	21	42**
Total aerobic mesophilic bacteria	PS	4.50 \pm 0.03 ^c	9.91 \pm 0.07	7.72 \pm 0.07 ^a
	DRD	6.08 \pm 0.17 ^b	9.58 \pm 0.40	7.13 \pm 0.25 ^b
	PS+DRD	6.89 \pm 0.08 ^a	9.96 \pm 0.04	7.82 \pm 0.04 ^a
<i>Salmonella-Shigella</i>	PS	-	3.80 \pm 0.03	-
	DRD	-	3.85 \pm 0.03	-
	PS+DRD	-	3.88 \pm 0.05	-
<i>Enterobacteriaceae</i>	PS	-	5.72 \pm 0.13	1.34 \pm 0.18 ^a
	DRD	-	5.48 \pm 0.09	0.56 \pm 0.31 ^b
	PS+DRD	-	5.90 \pm 0.11	1.46 \pm 0.09 ^a
<i>Coliforms</i>	PS	-	4.04 \pm 0.10	-
	DRD	-	3.97 \pm 0.14	-
	PS+DRD	-	3.94 \pm 0.02	-
<i>E. coli</i>	PS	-	3.08 \pm 0.07	-
	DRD	-	2.97 \pm 0.14	-
	PS+DRD	-	2.83 \pm 0.09	-
Enterococci	PS	0	4.69 \pm 0.15	7.39 \pm 0.10 ^a
	DRD	3.56 \pm 0.06 ^a	4.43 \pm 0.17	6.54 \pm 0.20 ^b
	PS+DRD	3.34 \pm 0.06 ^a	4.59 \pm 0.12	7.42 \pm 0.24 ^a
<i>S. aureus</i>	PS	-	5.46 \pm 0.09	0.99 \pm 0.08 ^a
	DRD	-	5.49 \pm 0.05	0.31 \pm 0.31 ^b
	PS+DRD	-	5.35 \pm 0.09	1.29 \pm 0.04 ^a
Yeasts and moulds	PS	3.20 \pm 0.09 ^b	7.69 \pm 0.16	7.26 \pm 0.30
	DRD	5.48 \pm 0.05 ^a	7.52 \pm 0.07	6.29 \pm 0.32
	PS+DRD	5.31 \pm 0.03 ^a	7.30 \pm 0.21	6.84 \pm 0.07

* $P < 0.001$, ** $P < 0.05$

^{a-c} Within columns, within litter materials, means with common superscripts do not differ ($P > 0.05$)

In Figure 1 the weekly pH values of litter materials are presented. While initially all the litter material groups were acidic, it was found that both the DRD and PS+DRD groups were acidic at day 21, but only the DRD group was acidic at day 42.

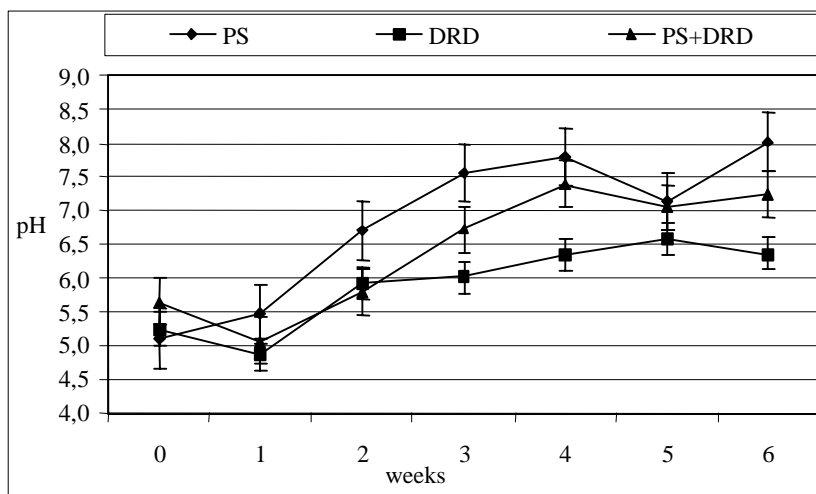


Figure 1 Weekly pH levels (with s.e. bars) of different litter material in the pens of boilers up to an age of 42 days

Conclusion

It was observed that the use of DRD as a litter material for broilers resulted in a decrease in the occurrence of pathogen microorganisms especially at the last stages of the finishing period. It is highly likely that this might have resulted from both the essential oils content in DRD and the effect of a decrease in the pH level of litter towards the last stages of the finishing period. These results showed that DRD as an industrial waste material in Turkey could be recycled and used as an enriched organic fertilizer in crop production, as well as decreasing the amount of pathogenic microorganisms in poultry production. Moreover, the essence of roses accumulated in the poultry houses provided less of a nuisance smell in the environment for workers than the wood shavings. The present study showed that as an alternative litter material, DRD caused antimicrobial and antibacterial effects during broiler production. However, further research with inoculated litters and for common use with not only DRD but also rose and other herbal species extracts are required to clarify the antimicrobial and antibacterial effects of the essential oils.

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