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Diallel Analysis on Breast and Thigh Muscle Traits in the Cross of Three South African Indigenous Chicken Genotypes

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ABSTRACT

The present study aimed to estimate carcass characteristics of pure and crossbred chickens produced from three parental populations. A 3 × 3 complete diallel mating system involving three indigenous breeds, namely Potchefstroom Koekoek (P), Venda (V), and Ovambo (O), was used to produce three purebred (P × P, V × V, O × O), three crossbreds (P × O, P × V, O × V) and three reciprocals (O × P, V × P, V × O). The nine genetic groups were reared from hatch to 10 weeks of age in an open house with deep litter. At 10 weeks of age, six chickens per genetic group were randomly selected for slaughter. After slaughtering the breast and thigh muscles samples for analysis of the carcass characteristics (Meat colour, meat pH, and Shear force). The results showed that the Potchefstroom Koekoek breed had higher values in all colour indicators, L* (lightness), a* (redness), and b* (yellowness), compared to the other chicken breeds. The Potchefstroom Koekoek and P × O breed had higher pH values ranging from 5.66 to 6 at two hours post-slaughter and from 5.54 to 6.38 at 24 hours post-slaughter. The pH declines in all the nine genetic groups after two to 24 hours, with the exception of the crossbred P × O, which increased from 6.06 to 6.38. In terms of shear force, the O × P had the highest shear value, ranging from 35.89N to 74.80N, compared to other genetic groups. Potchefstroom Koekoek had normal meat colour and pH, whereas the Venda breed had tougher meat than other genotypes. The results of the present study might be useful for local chicken farmers to improve carcass traits.

Keywords: Crossbred, Meat colour, Meat pH, Purebred, Shear force.

INTRODUCTION

Diallel cross is the crossbreeding among several genotypes and data analysis from such crosses (Khalil and Khadiga, 2020). Crossbreeding is one of the tools that can be used in genetic variation (Wang et al., 2019). This crossbreeding which is the mating of two individuals with different breed compositions, is one type of a larger class of mating systems known as out-breeding (Khalil and Khadiga, 2020). There are many South African indigenous chicken genotypes, including Potchefstroom Koekoek, Venda, and Ovamba chicken (Manyelo et al., 2020). The Potchefstroom Koekoek chicken genotype, developed by crossing Black Australorp cockerels with White Leghorn hens and Plymouth Rock, reaches sexual maturity after 130 days and is mainly categorized by black and white speckled colour patterns (Mutibvu et al., 2020). The Venda chicken genotype, discovered in the Venda region of South Africa, is a dual-purpose breed with multicoloured patterns predominated by black and white colours (Manyelo et al., 2020). On the other hand, the Ovambo chicken genotype, first discovered in the Ovamboland of northern Namibia, is predominantly dark to black in colour and can reach sexual maturity after 140 days (Mutibvu et al., 2020). Tyasi et al. (2019) conducted a diallel cross of three South African indigenous chicken genotypes, namely chicken genotypes Potchefstroom Koekoek, Venda, and Ovamba, to investigate the crossbreeding effects on growth traits. Thus far, there had been limited studies of the crossbreeding effects on breast and thigh muscles in Potchefstroom Koekoek, Ovambo, and Venda chicken breeds. Hence, the objective of the study was to determine the effect of crossbreeding on meat pH, meat colour and meat tenderness on breast and thigh muscle traits. The current study will aid indigenous chicken farmers to improve carcass characteristics through crossbreeding.

MATERIALS AND METHODS

Ethical approval

This research was carried out in accordance with the standard operation procedures of the Animal Research and Ethics Committee (AREC) at the University of Limpopo, South Africa.

Study area

The present study was conducted on the experimental farm of the University of Limpopo in the Limpopo Province of South Africa. The farm is located about 10 kilometers northwest of the university. The mean temperatures in winter (April to July) range between 10.1 and 28.4 °C and in summer (August to March) between 18 and 36 °C. The annual rainfall ranges between 446.8 and 468.4 mm.

Preparation of the house

The house was prepared as explained in Alabi et al. (2020). Briefly, the incubator and all the equipment such as drinkers, feeders, and wire separators were thoroughly cleaned and disinfected.

Chickens management

Ten hens of each breed were assigned randomly to be mated with two roosters of each breed. The eggs were collected daily by breeds and crossbreds and then hatched according to breeds and crossbreds. separately Accordingly, nine genetic groups of $P \times P$, $V \times V$, $O \times O$, $P \times V$, $P \times N$, $V \times P$, $V \times O$, $O \times P$, and $O \times V$ chickens were obtained. The hatched chickens were wing banded until eight weeks of age, followed by leg bands to keep their breed and crossbred group identity. The chickens were kept together on a litter floor in a semi-open house, which was partitioned according to their breeds and crossbreds. They were medicated in a similar way and were subjected to the same management, hygiene, and climatic conditions. During the brooding and rearing periods, all chickens were fed ad libitum using a standard commercial starter (21 % Crude Protein (CP) and 3,000 kcal Metabolizable Energy (ME)/kg) from hatching time until four weeks of age, followed by a grower diet (18 % CP and 2,900 kcal ME/kg) up to ten weeks of age. Water was provided ad libitum. Artificial heat (32 °C) using infrared lights and a continuous light program was provided. The ventilation was controlled using curtain rails.

Meat pH measurements

A portable pH meter with a fibre-optic probe (CRISON pH 25 Instruments S.A., Alella, Spain) was used to measure the pH (pH 2 and 24 hours post mortem) of the carcasses. The pHu was first calibrated using standard solutions of pH 4, pH 7, and pH 9 (CRISON Instruments, SA, and Spain). The measurements were then performed with a sharpened metal sheath to prevent probe breakage due to the raw meat contamination.

Determination of meat colour

The colour of the meat (L*: Lightness, a*: Redness, and b*: Yellowness) was determined two hours after slaughter (Commission International De I' Eclairage, 1976). A refrigerated portable vehicle was used to move the carcass from the slaughtering place to the University of Limpopo, Department of Animal Production laboratory one hour after the slaughter. A Minolta colour-guide 45/0 BYK-Gardener GmbH machine with a 20mm diameter measurement and an illuminant D65-daylight, from 100 standard observers was used for the colour measurement. The machine was calibrated each time before measurements were taken using the standard green, black and white colour samples provided for this purpose. The readings were taken by rotating the Colour Guide 900 between measurements to obtain the average value for the colour.

Determination of meat tenderness

The tenderness of the chicken was determined using the Instron- Warner-Bratzler Shear Force (WBSF). After cooking, subsamples of specified core diameter were cored parallel to the grain of the meat. Three subsamples with a core diameter of 10 mm were cored parallel to the grain of the meat. The samples were sheared perpendicular to the fiber direction using a Warner Bratzler (WB) shear device mounted on an Instron (Model 3344) Universal Testing apparatus (the crosshead speed at 400mm/minutes, one shear of each core). The mean maximum load (N) was recorded for the batch.

Data analysis

The Generalized Linear Models procedure (PROC GLM) of Statistical Analysis System (SAS, 2019), version 9.4, was used to determine the effect of the chicken genotype on the chicken quality attributes (pH, L*, a*, b* and WBSF values).

$$y_i = \mu + \alpha i + \varepsilon i$$

Where,

Vi = Response variable (pH, L, a*, b* and WBSF), μ is the overall mean, α i denoted breed effect (V × V, O × O, P × P, V × O, V × P, O × V, O × P, P × V and P × O genotypes), and ϵ i refers to random error term. Differences considered significant (p \leq 0.05) were compared with Fisher's protected Least Significant Difference (LSD test).

RESULTS

The meat colour of different genotypes for the breast muscle is presented in Table 1. The results showed a significant difference (p < 0.05) in the colour of the breast muscle between the nine genetic groups. In terms of lightness, the Ovambo chicken genotype had a higher value, followed by $O \times V$ chicken genotype. However, there were no statistically significant differences (p > 0.05)between crosses ($P \times O$, $P \times V$, and $O \times V$) and reciprocals $(V \times P, V \times O, and O \times P)$. For redness, the Potchefstroom Koekoek genotype had a higher value (p < 0.05) followed by the $P \times O$ chicken genotype. The results also indicated that $O \times V$ had statistically insignificant differences (p > 0.05) with reciprocals (V \times P, V \times O, and O \times P) as well as $V \times V$ and $P \times V$. In yellowness, the Potchefstroom Koekoek chicken genotype also had a higher value (p < p0.05) than the other chicken genotypes. The results showed that there were no statistically significant differences (p > 0.05) between V \times V and crosses (P \times O, $P \times V$, and $O \times V$). These results also indicated that there were no remarkable differences (p > 0.05) between V \times V and the reciprocal values ($O \times P, V \times P$).

Table 2 shows a meat colour of different genotypes for the thigh muscle. The results showed a significant difference (p < 0.05) in thigh muscle colour between the nine genetic groups. In terms of lightness, Potchefstroom Koekoek genotype, and Venda chicken genotype had a higher value (p < 0.05), followed by Ovambo chicken genotype and the crosses (P \times O, P \times V, and O \times V) chicken genotypes. However, no statistically significant differences (p > 0.05) between reciprocals (V \times P, V \times O, $O \times P$) were observed. In redness, The findings recognized that no statistically significant differences (p > 0.05) were observed between $P \times P$, $O \times O$ purebreds, $P \times O$, $P \times V$, $O \times V$ crosses, and $V \times O$ reciprocal. In yellowness, the Potchefstroom Koekoek chicken genotype had a higher value (p < 0.05) than the other chicken genotypes. The results also showed that there were no statistically significant differences (p > 0.05) between $O \times O$, $V \times V$, and $O \times V$ cross and $O \times P$ reciprocal.

The meat pH values of breast muscle at 2 and 24 hours post-mortem in purebreds, crossbreds, and

reciprocals are presented in Figure 1. At a pH of 2 hours post-mortem, the findings indicated that the cross of $P \times O$ had a higher pH value (5.70 ± 0.09) (p < 0.05) than the other genetic groups. The results indicated that no statistically significant differences (p > 0.05) between purebreds' $P \times P$ (5.66 ± 0.01), $O \times O$ (5.61 ± 0.02), and V \times V (5.64 \pm 0.03) were observed. The results also recognized that there were no remarkable differences (p > p)0.05) between the reciprocal genetic groups O \times P (5.57 \pm 0.16), V × P (5.52 \pm 0.02), and V × O (5.55 \pm 0.01) were observed. At a pH of 24 hours post-mortem, the results indicated that the Potchefstroom Koekoek chicken genotype had a higher pH value (5.64 ± 0.03) (p < 0.05) than the other chicken genotypes. The results indicated that there were no statistically significant differences (p > p)0.05) between reciprocal genetic groups O \times P (5.47 \pm 0.04), V × P (5.53 \pm 0.02), and V × O (5.47 \pm 0.04), were observed.Meat pH of thigh muscle at 2 and 24 hours' postmortem in purebreds, crossbreds, and reciprocals is presented in Figure 2. At a pH of 2 hours post-mortem, the results showed that the cross of $P \times O$ had a higher pH value (6.06 \pm 0.07) (p < 0.05) than the other genetic groups. The results indicated that there were no remarkable differences (p > 0.05) between purebreds' P \times P (6.00 \pm 0.04), and V \times V (5.98 \pm 0.08), and reciprocal V \times P (5.96 ± 0.06) chicken genotype were observed. At a pH of 24 hours post-mortem, the findings recognized that the cross of $P \times O$ chicken genotype had a higher pH value (5.70 ± 0.09) (p < 0.05) than the other chicken genotypes. There were no statistically significant differences (p > p)0.05) between V \times V (5.98 \pm 0.08), cross O \times V (5.83 \pm 0.07), and reciprocals V \times P (5.96 \pm 0.06) and V \times O (5.90 \pm 0.03) were observed.

Table 3 shows the results of the shear force for thigh and breast muscles. In purebreds, the results showed that P \times P had a higher shear force value as compared with other purebreds but there were no statistically significant differences (p > 0.05) between P \times P and O \times O in thigh muscle. Whereas in breast muscle, there were no statistically significant differences (p > 0.05) observed between purebreds. In crosses, the findings indicated that there were no statistically significant differences among all crosses (p > 0.05). While in breast muscle, $O \times V$ had a higher shear force value (p < 0.05) followed by $O \times V$, respectively. With respect to reciprocals, $V \times O$ had a higher shear force value (p < 0.05) but no statistically significant differences (p > 0.05) observed between $O \times P$ and $V \times P$. However, in breast muscle, there were no statistically significant differences (p > 0.05) detected between $O \times P$ and $V \times O$, respectively.

	Meat colour indicators		L *	a*	b*
Genotypes					
		$\mathbf{P} \times \mathbf{P}$	43.56 ± 0.02^{d}	9.39 ± 0.51^{a}	10.68 ± 0.88^{a}
Purebreds		$\mathbf{O} \times \mathbf{O}$	56.54 ± 1.50^a	1.68 ± 0.30^{e}	6.69 ± 0.61^{c}
		$\mathbf{V}\times\mathbf{V}$	44.79 ± 0.89^{cd}	5.13 ± 0.81^{cd}	9.15 ± 0.77^{abc}
		$\mathbf{P} \times \mathbf{O}$	48.03 ± 3.09^{bcd}	6.51 ± 0.43^{b}	8.96 ± 0.90^{abc}
Crosses		$\mathbf{P}\times\mathbf{V}$	47.41 ± 0.38^{bc}	6.43 ± 0.29^{bc}	8.99 ± 0.84^{abc}
		$\mathbf{O}\times\mathbf{V}$	50.93 ± 2.45^{b}	4.90 ± 0.29^{cd}	9.36 ± 1.11^{ab}
		$\mathbf{O} \times \mathbf{P}$	48.84 ± 2.35^{bc}	5.05 ± 0.65^{cd}	7.44 ± 1.22^{bcd}
Reciprocals		$\boldsymbol{V}\times\boldsymbol{P}$	$46.93 \pm 0.55^{\circ}$	4.02 ± 0.38^{d}	7.60 ± 1.22^{bcd}
		$\mathbf{V} imes \mathbf{O}$	47.64 ± 0.11^{bcd}	3.75 ± 0.47^{d}	5.13 ± 1.11^{d}

Table 1. Meat colour of breast muscle in purebreds, crosses, and reciprocals of three South African indigenous chicken genotypes (Potchefstroom Koekoek, Ovambo, and Venda)

^{a-d}: Means in the same column with the same letters did not differ significantly (p > 0.05), $P \times P$: Potchefstroom Koekoek, $O \times O$: Ovambo, $V \times V$: Venda, L*: Lightness, a*: Redness, b*: Yellowness.

Table 2. Meat colour of thigh muscle in purebreds, crosses,	and reciprocals of three South African indigenous chicken
genotypes (Potchefstroom Koekoek, Ovambo, and Venda).	

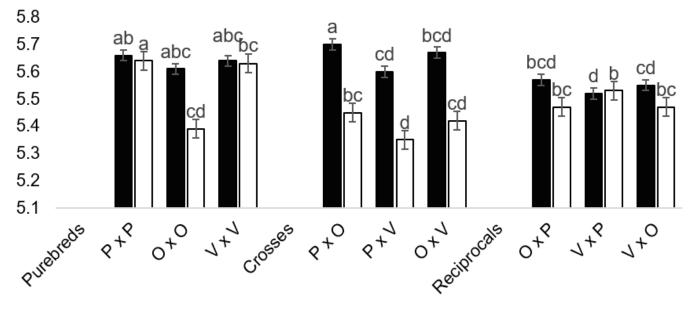
	Meat colour indicators		L *	a*	b*
Genotypes			L.	a.	D.
		$\mathbf{P} \times \mathbf{P}$	52.09 ± 1.12^{a}	$8.95\pm0.58^{\rm a}$	12.87 ± 1.16^{a}
Purebreds		$\mathbf{O} \times \mathbf{O}$	46.29 ± 0.59^b	8.54 ± 0.53^a	9.26 ± 0.31^{b}
		$\mathbf{V}\times\mathbf{V}$	50.83 ± 0.36^{a}	6.47 ± 0.53^{c}	7.36 ± 1.59^{bc}
		$\mathbf{P} \times \mathbf{O}$	45.92 ± 0.93^{b}	$8.06\pm1.15^{\rm a}$	$6.56\pm2.08^{\rm c}$
Crosses		$\mathbf{P}\times\mathbf{V}$	43.11 ± 0.74^{b}	$8.74\pm0.62^{\rm a}$	4.53 ± 0.78^{d}
		$\mathbf{O}\times\mathbf{V}$	$45.67 \pm 1.11^{\text{b}}$	8.00 ± 1.37^{a}	7.77 ± 2.04^{bc}
		$\mathbf{O} \times \mathbf{P}$	46.80 ± 0.71^{b}	7.43 ± 0.36^{b}	7.03 ± 0.77^{bc}
Reciprocals		$\boldsymbol{V}\times\boldsymbol{P}$	46.60 ± 0.82^{b}	6.78 ± 0.11^{bc}	$6.81 \pm 1.40^{\text{c}}$
		$\mathbf{V}\times\mathbf{O}$	47.55 ± 0.45^{b}	$8.30\pm0.32^{\rm a}$	$6.27 \pm 1.62^{\circ}$

^{a-d} Means in the same column with the same letters did not significantly differ (p > 0.05), $P \times P$: Potchefstroom Koekoek, $O \times O$: Ovambo, $V \times V$: Venda, L*: Lightness, a*: Redness, b*: Yellowness.

Table 3. Comparison of shear force in purebreds, cros	es, and reciprocal	s of three	South African	indigenous c	chicken
genotypes (Potchefstroom Koekoek, Ovambo, and Venda).					

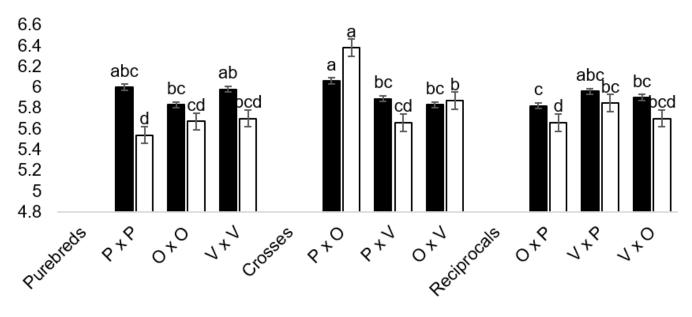
Genotypes	Shear force	Thigh WBSF	Breast WBSF
	$P \times P$	44.15 ± 7.45^a	$58.82\pm5.92^{\rm c}$
Purebreds	$O \times O$	40.50 ± 6.78^{ab}	51.94 ± 3.87^{cd}
	$V \times V$	32.50 ± 1.55^{bc}	59.07 ± 2.81^{c}
	P× O	31.38 ± 2.04^{bc}	$55.51 \pm 9.64^{\circ}$
Crosses	$P \times V$	29.25 ± 2.01^{c}	43.62 ± 0.64^d
	$O \times V$	31.73 ± 3.05^{bc}	$61.23\pm4.36^{\text{b}}$
	$O \times P$	$35.89 \pm 1.36^{\text{b}}$	74.80 ± 7.15^a
Reciprocals	$V \times P$	42.33 ± 3.56^{ab}	65.41 ± 6.35^b
	$V \times O$	46.09 ± 6.36^{a}	72.34 ± 8.26^{a}

^{a-d} Means with the same column with the same letters did not significantly differ (p>0.05), $P \times P$: Potchefstroom Koekoek, $O \times O$: Ovambo, $V \times V$: Venda. WBSF: Warner Braztler Shear Force



■pH2h □pH24h

Figure 1. Meat pH of breast muscle 2 and 24 hours in purebreds, crosses, and reciprocals of three South African indigenous chicken genotypes. ^{a-d}: Means that the same column with the same letters did not significantly differ (p > 0.05), $P \times P$: Potchefstroom Koekoek, $O \times O$: Ovambo, $V \times V$: Venda, $pH_{2h} = pH 2$ hours post-mortem, $pH_{24h} = pH 24$ hours post-mortem.



∎pH2h □pH24h

Figure 2. Meat pH of thigh muscle at 2 and 24 hours in purebreds, crosses, and reciprocals of three South African indigenous chicken genotypes. ^{a-d}: Means with the same column with the same letters did not significantly differ (p>0.05), $P \times P$: Potchefstroom Koekoek, $O \times O$: Ovambo, $V \times V$: Venda, $pH_{2h} = pH 2$ hours' post-mortem, $pH_{24h} = pH 24$ hours' post-mortem.

DISCUSSION

Crossbreeding may possibly be used to achieve genetic improvement with or without genetic selection in the parental lines (Wang et al., 2019). The influence of crossbreeding on meat color, pH, and tenderness in breast and thigh muscles of nine genetic groups namely; purebreds (P \times P, O \times O, V \times V), crossbreds (P \times O, P \times V, $O \times V$) and reciprocals ($O \times P$, $V \times P$, $V \times O$) were explored in this study. Meat colour is one of the factors which the consumer uses as a measure of acceptance. For example, a pink interior is considered desirable in a processed whereas a pink or reddish colour is highly undesirable in fully cooked poultry meat (Singh et al., 2021). In the present study, meat color findings suggested that the breast and thigh of the Ovambo chicken genotype were paler (high L*), more yellow (high b*) and less red (low a*) than other genetic groups. Potchefstroom Koekoek chicken genotype had higher values in all color indicators, L* (lightness), a* (redness), and b* (yellowness) as compared to the other chicken breeds. It was however observed that the Venda breed had a high L* (lightness) which was normal between 43 and 53. The pH of chicken meat for the current study was a significant source of variability amongst the nine genetic groups at different hours (2 hours, 24 hours) after slaughter. The pH declined in all the nine genetic groups at 2 to 24 hours with the exception of crossbred, $P \times O$ which increased from 6.06 to 6.38. The tenderness of the meat was also examined with a Warner-Bratzler shear force. The Warner-Bratzler shear force is an objective measure of tenderness used in the research laboratory to evaluate relative differences in tenderness or toughness of meat products (Wang et al., 2019). The results of the present study indicated that shear force values of breast muscle were higher than those of the thigh muscle. Similar results were also reported by Wang et al. (2019). Singh et al. (2021) compared the carcass characteristics of a slowgrowing crossbred broiler to a fast-growing broiler and found that the pH at 24-hour post-mortem was 5.88 for a fast-growing broiler and 5.74 for a slow-growing broiler. The breast color of the fast-growing broiler had a lightness (L*) of 51.93, a redness (a*) of 208, and a yellowness (b*) of 4.97, while the slow-growing broiler had an (L*) of 45.39, an (a*) of 2.64, and a (b*) of 3.84, respectively. Mueller et al. (2020) studied the meat quality of two dualpurpose and one-layer hybrid chickens for 67 or 84 days compared to a slow-growing broiler. The results showed that the pH in the breast muscle ranged from 5.00 to 6.00 at 24 hours post mortem, while the breast colour ranged from 47.0 to 59.0 for L*, 1.11 to 3.33 for a*, and -0.22 to 1.06 for b*. Devatkal et al. (2019) recognized that the breast shear force of the slow-growing broiler was 11.09 and 12.47 for the fast-growing broiler. Mueller et al. (2018) indicated that the breast meat pH at 24-hour postmortem for Ross PM3 chicken was 6.25, Sasso S1 chicken 5.92, Lohmann Dual chicken 5.82, Belgian Malines chicken 5.91, Schweizerhuhu chicken 5.73, and Lohmann Brown Plus chicken 5.90, while the breast colour ranged from L* 49.0 to 54.6, a* 1.26 to 3.58, and b* 0.09 to 0.72, respectively. These studies are inconsistent with the current study and the variations might be due to different genotypes, age, nutrition, and environment. Tyasi et al. (2019) discovered that the Potchefstroom Koekoek chicken genotype had a positive influence on the general ability to combine and the crossbred between the Potchefstroom Koekoek and the Venda chicken genotype had a positive influence on the specific ability to combine, and thus suggested that the Potchefstroom Koekoek chicken genotype could possibly be used for crossbreeding with other indigenous chicken genotypes to improve growth traits.

CONCLUSION

Crossbreeding had a remarkable effect on meat colour, pH, and tenderness in a diallel cross of the Potchefstroom Koekoek, Ovambo, and Venda chicken genotypes. The Potchefstroom Koekoek genotype had normal meat colour and pH whereas the Venda chicken genotype had tougher meat than other chicken genotypes. It is recommended that crossbreeding could be introduced to improve genetic effects on carcass quality traits.

DECLARATION

Acknowledgments

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Author's contributions

Thobela L Tyasi conducted the experiment, analyzed the data and wrote the manuscript. Jones W Ng'ambi and David Norris designed and oversaw the running of the experiment.

Competing interests

The authors declare that they have no conflict of interest.

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