



Immune and Growth Response of Indigenous Pedi Goats Vaccinated with Blanthrax to an Inclusion of *Moringa oleifera* (Drumstick Tree) in *Cenchrus ciliaris* (Buffel Grass) Hay-Based Diet

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Abstract | The study aimed to ascertain the immunomodulatory effects of *Moringa oleifera* (*M. oleifera*) leaves supplemented to the diets of wether BaPedi goats following vaccination with blanthrax vaccine. Twelve clinically healthy BaPedi goats with an average body weight of 19 ± 1.47 kg and an average age of 11 ± 0.26 months were randomly selected from the flock at the University of Limpopo experimental farm. The experiment was conducted in three phases which are adaption, vaccination and moringa inclusion over 42 days. At the end of the first week of the trial, all the experimental goats were vaccinated with 2 millilitres of blanthrax vaccine per goat via the subcutaneous route. Three experimental diets were formulated by replacing a conventional supplement of Lucerne with *M. oleifera*. The inclusion levels varied from 0% to 50%. Data on the growth and haematological parameters of the animals used in the study were analysed using a general linear model (GLM) procedure in a completely randomized design. The results indicated that only platelet counts, monocyte counts and mean corpuscular volume (MCV) showed significant differences ($p < 0.05$) amongst the 11 blood profiles observed in this study. There were significant differences ($p < 0.05$) in body weight gain (BWG), growth rate (GR) and metabolic weight gain (MWG). No significant differences ($p > 0.05$) were observed for feed intake (FI) and feed conversion ratio (FCR). Results of the present study suggest that *M. oleifera* leaves can be used as a feed supplement at 20% and 50% inclusion levels without having any adverse effects on blood parameters and growth performance.

Keywords | Dietary supplementation, Growth performance, Haematology, Natural forages, Vaccination

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INTRODUCTION

Goat production plays an integral role in the world especially in communal areas because they can survive under tough climatic conditions (Fikru and Omer, 2015). The indigenous goats reared in communal areas are commonly perceived as less productive than exotic breeds because successful production in communal areas is often affected by various challenges such as malnutrition, poor health, and welfare-related issues caused by the unavailability of resources (Gwanzura et al., 2011; Abreu et al., 2017). The farmers in communal areas rarely conduct

livestock management practices such as vaccinations, moreover, disease cases in animals following vaccination have been reported (Monau et al., 2020). Vaccine negative effects in some cases could be due to underlining conditions such as malnutrition and inappropriate management practices, especially in drought-prone areas (Ayele et al., 2016; Kniffen and Comerford, 2021). An example of a vaccine that is commonly used by farmers in the Limpopo province is blanthrax. It is a combined vaccine that is commonly used worldwide because it is perceived to provide good protection against anthrax and blackleg (Wilson, 2003; Ndeereh et al., 2012).

Although trees and shrubs have been used as a source of nutrients and for their medicinal benefits for livestock, most of these plants do not contain optimal nutrients and components required for ideal performance (Ogbe and Affiku, 2011; Nouman et al., 2014). This often leads to poor immune responses following vaccination. Documentation of the use of plants with herbal and nutritional properties is essential, particularly given the escalating costs of drugs and feed as well as the escalating resistance of pathogens to drugs (Zhang and Ma, 2018).

Interest has developed in the *Moringa oleifera* (*M. oleifera*) tree because of its fast growth, higher nutritional traits, medicinal properties, and utilization as a livestock fodder crop (Verma et al., 2009; Nouman et al., 2014). *Moringa oleifera* thrives best in the tropics and subtropical areas such as eastern and southern Africa (Moyo et al., 2012; Nouman et al., 2014). The levels of harmful substances such as tannins and phytates are very low. Therefore, it is recommended that *M. oleifera* can be used for its health benefits in livestock (Udom and Idiong, 2011; Nouman et al., 2014). Several studies explored the nutritional and medicinal properties of *M. oleifera* and found it to be beneficial as a supplement in livestock feeds (Ogbe and Affiku, 2011; Moyo et al., 2012; Nouman et al., 2014). The dried leaves of *M. oleifera* have average crude protein (CP) levels of up to 30.3%, it is rich in vitamins and minerals needed by the animal's body to create immune cells for a strong defence system (Moyo et al., 2011; Adouko et al., 2020). Some studies reported trace levels of non-nutritive elements found in *M. oleifera* leaves, such as Barium (Ba), Cadmium (Cd) and Argentum (Ag) which could have arisen from environmental contaminations (Pakade et al., 2013; Biel et al., 2016). These variations may be due to factors such as the different geographic locations, soil type, stage of growth of the tree, the season of harvesting, and storage conditions before analyses (Mulyaningsih and Yusuf, 2018).

Moringa contains antioxidants such as polyphenols in acceptable ranges that may aid in boosting the immune profile of the animal (Ma et al., 2018). Therefore, this study aimed to ascertain the immunomodulatory and growth effects of *M. oleifera* supplementation to the diets of castrated indigenous BaPedi goats following vaccination with the blanthrax vaccine. The findings of the present study will aid in decision making on the utilization of *M. oleifera* as a feed supplement in communal goat production.

MATERIALS AND METHODS

All procedures used in this experiment followed the ethical standards of the University of South Africa (2019_CAES_AREC_131) and the University of Limpopo (AREC/10/2019:1R) Animal Ethics Committees. The

experiment proceeded after both ethical clearances were obtained.

The study was conducted at the University of Limpopo experimental farm in the summer season of 2020 (October-December). The farm is situated in the tropic of Capricorn zone, in the southern region of the Limpopo Province. The average temperatures range from 20 to 36 °C during the wet season (November to January) and between 5 and 25 °C during the dry season (May to July) (Brown et al., 2016). Steel, individual well-ventilated pens, with an area of 3.75 m² were installed at the site for the experiment. The pens were equipped with a feeder and a drinker. The experimental site was thoroughly cleaned and disinfected daily to avoid a build-up of infections such as pneumonia.

Twelve clinically healthy BaPedi goats with an average of 19 ± 1.47 kg body weight and 11±0.26 months of age were randomly selected from the flock at the University of Limpopo experimental farm (Lee et al., 2014; Brown et al., 2016b). Only male goats were used in this study because the sex of the animal can affect haematological values in farm animals (Binta et al., 1996; Tibbo et al., 2004). Health assessments and clinical observations (body temperature, respiratory rate, pulse rate, heart rate, signs of gastrointestinal malfunction such as diarrhoea/constipation, examination of mucous membranes of the conjunctiva to evaluate the circulatory system and signs of anaemia) were done daily throughout data collection to ensure only healthy goats were used. The goats were randomly divided and allocated a treatment diet in a completely randomized design using an individual goat as a replicate (Steel and Torrie, 1980). From the first week until the fourth week of the trial, the goats were fed a control basal diet comprising 90.77% Lucerne (*Medicago sativa*) and 9.23% Buffel grass (*Cenchrus ciliaris*) for acclimatization of their rumen microbes (Odenyo et al., 1997). On the fifth and sixth weeks, the goats were fed diets comprising of varying levels of *M. oleifera* (Table 1). The animals had *ad-libitum* access to fresh water throughout the data collection period. At the end of the first week of the trial, all the experimental goats were vaccinated with 2 millilitres of blanthrax vaccine per goat via the subcutaneous route as prescribed by the vaccine manufacturer (Intervet South Africa, (Pty) Ltd). The objective of vaccinating the goats with a blanthrax vaccine was to evaluate the immune response following vaccination and compare it with the immune response following supplementation of the experimental diet. One of the goats was removed because it had displayed signs of dietary intolerances in the first week of the trial. Thus, the number of experimental units and animals used for the data analysis was eleven (three animals for the T₁C_{9.23}L_{90.77}M₀ diet) (Magasa and Mbassa, 1988).

Table 1: Formulation of dietary treatments used for the study.

Treatment code	Treatment description	Number of animals
T ₁ C _{9.23} L _{90.77} M ₀	Wether goats were fed <i>ad-libitum</i> a basal diet, 9.23 % <i>C. ciliaris</i> grass hay, and 90.77% Lucerne hay without <i>M. oleifera</i> .	3
T ₂ C ₆₄ L ₁₆ M ₂₀	Wether goats were fed <i>ad-libitum</i> 64 % <i>C. ciliaris</i> grass hay and 16 % percent Lucerne hay containing 20 % of <i>M. oleifera</i> .	4
T ₃ C ₅₀ L ₀ M ₅₀	Wether goats were fed <i>ad-libitum</i> 50 % <i>C. ciliaris</i> grass hay and 0 % percent Lucerne hay containing 50 % of <i>M. oleifera</i> .	4

L: Lucerne hay; C: *C. ciliaris* grass hay; M: *M. oleifera*

Fresh *M. oleifera* leaves and *Medicago sativa* (*M. sativa*) were harvested and air-dried on a concrete floor at room temperature for 96 hours and turned several times until they were crispy in touch, while retaining their greenish colouration. The dried leaves of *M. oleifera* and Lucerne were then stored at room temperature until needed for inclusion in the experimental diets. *Cenchrus ciliaris* (*C. ciliaris*) which is one of the palatable and perennial grass species that are native to the Limpopo province was used in this study (Smit, 2005). It contains an average of 6.1 % CP. Lucerne hay was used to supplement the insufficient protein content of Buffel grass. Lucerne contains an average of 11.4% CP. Three experimental diets were formulated by replacing a conventional supplement of Lucerne with *M. oleifera*. The inclusion levels varied from 0% to 50% as presented in Table 1 (Babeker and Bdalbagi, 2015; Su and Chen, 2020). *Moringa oleifera* leaves utilised in the present study contained traces of non-nutritive elements, such as Ba, Cd, and Ag which could have arisen from environmental contaminations (Pakade et al., 2013). However, there were no signs of toxicity observed in the animals due to the presence of heavy metal trace elements in the leaves. This could be attributed to the low levels of their presence (Pakade et al., 2013; Biel et al., 2016).

Goats were fed experimental diets that were 4% of their body weight daily. Analyses of CP, ash and NDF in dietary treatments were done at the Animal Nutrition Laboratory, University of Limpopo as described by AOAC (1990) (Table 2). The mineral composition was determined at the Limpopo Agro-Food Technology Station (LATS) Laboratory, University of Limpopo as described by ISO 11885 (2007).

The experiment was conducted in three phases for 42 days. The first phase was the adaptation phase which lasted for 7 days. The second phase was the vaccination phase, which lasted for 21 days. The duration of this phase depended on the establishment of immunity in goats which according to the manufacturer takes a minimum of 14 days. The third phase which lasted for 14 days was the inclusion phase of varying levels of *M. oleifera*.

Five millilitres of blood was collected from each animal

via the external jugular vein on day 0 and at the end of each phase. Blood parameter results for day 0 were used as baseline values, this was done to evaluate changes in the haematological responses of the goats (Brown et al., 2016b). Vacutainer tubes containing ethylenediaminetetraacetic acid (EDTA) as an anticoagulant were used for blood collection (Daramola et al., 2005; Aikhuomobhogbe and Orheruata, 2006; Brown et al., 2016b). The blood samples were stored in a cooler box containing ice packs and they were immediately transported to the University of Pretoria, Department of Companion Animal Clinical Studies, Gauteng Province for haematological analyses. All blood parameter results were referenced with day 0 as baseline values according to Brown et al. (2016).

Table 2: Nutrient composition of experimental diets used in the study.

Nutrients	T ₁ C _{9.23} L _{90.77} M ₀	T ₂ C ₆₄ L ₁₆ M ₂₀	T ₃ C ₅₀ L ₀ M ₅₀
CP %	11.7	10.48	15
NDF %	52.53	50.36	39
Ash %	8.41	8.86	8.5
Zn (mg/kg)	11.81	2.25	4.25
Fe (mg/kg)	449.07	605.6	434.9
Mn (mg/kg)	69.57	129.21	98.78
Cu (mg/kg)	10.39	10.88	10.21
Mg (g/kg)	1.26	0.83	166.95
Na (g/kg)	1.22	0.52	23.55
P (g/kg)	2.87	2.06	73.4
Ca (g/kg)	8.48	3.41	523.45
K (g/kg)	13.82	13.35	550.2

T: Treatment; C: *C. ciliaris*; L: Lucerne; M: *M. oleifera*; CP: Crude protein; NDF: neutral detergent fibre; Zn: Zinc; Fe: Iron; Mn: Manganese; Cu: Copper; Mg: Magnesium; Na: Sodium; P: Phosphorus; Ca: Calcium; K: Potassium.

Bodyweight data was collected on day 0 from each goat and every seven days in the morning before feeding. The feed offered and refusals were weighed using an electronic scale each day from day 7 until day 42.

Feed intake (FI) was calculated as the difference between feed offered and feed refused. Live body weights were used

to calculate growth rate (GR) and feed conversion ratio (FCR). The metabolic weight gain (MWG) is expressed as the bodyweight gain (BWG) in grams (g) to the exponential of 0.75. The FCR was obtained by dividing feed intake by weight gain. The formulae used were:

$$FCR (kg) = \frac{FI}{BWG} \dots (1)$$

$$MWG (g) = BWG^{0.75} \dots (2)$$

$$GR = \frac{\text{Final body weight} - \text{initial body weight}}{\text{number of days}} \dots (3)$$

Descriptive statistics such as the mean, standard deviation, and standard error were calculated across different phases and treatments using Procedure of means (Proc mean) of Statistical Analysis Software version 9.4 (SAS, 2020). Data on blood parameters and growth performance was analysed using a general linear model (GLM) procedure. The following model was used:

$$Y_{ij} = \mu + T_i + e_{ij}$$

Where;

Y_{ij} = blood profiles or growth performance; μ = the overall mean; T_i = dietary treatments and e_{ij} = random error. Duncan's multiple range test was used for any significant differences ($p < 0.05$) amongst treatment means (Duncan, 1955).

RESULTS AND DISCUSSION

Measurements on blood parameters before and after vaccination were carried out during the study. There were no significant differences ($p > 0.05$) amongst means for the parameters except for MCV, platelet and monocyte counts ($p < 0.05$) (Table 3). Hb (80-120 g/dL), RBC (17-20 (*10)⁶ μ L), PCV (0.22-0.38 L/L), MCH (5.2-8.0 pg), MCHC (30-36 g/dL), WCC (6.8-20 (*10)³ μ L), monocytes (0-1%) and eosinophils (1-7%) for day 0 were within range of healthy goats (Jackson and Cockcroft, 2002; Daramola et al., 2005; Brown et al., 2016b). However, MCV (16-25 fL) and platelet counts (300- 600 (*10)⁹/L) for day 0 were out of range for healthy goats (Jackson and Cockcroft, 2002).

Table 3: Blood parameters across all phases from day 0- 42 and varying *M. oleifera* inclusion levels from day 29-42.

Parameters	Adaptation phase		Vaccination phase	Vaccination + <i>M. oleifera</i> phase		
	DAY 0 (Mean±SEM)*	DAY 7 (Mean±SEM)*	DAY 28 (Mean±SEM)*	VMO0	VMO20 DAY 42 (Mean±SEM)*	VMO50
Hb	92.91±3.30	92.67±5.16	84.64±4.08	79.33±2.40	90.00±4.45	83.50±8.38
RBC	17.83±0.70	17.42±0.85	16.40±0.89	15.57±0.88	16.92±0.68	16.41±1.88
HCT	0.28±0.01	0.28±0.02	0.26±0.01	0.23±0.01	0.28±0.02	0.25±0.03
MCV	15.64±0.22 ^{ab}	16.02±0.23 ^{ab}	15.62±0.20 ^{ab}	15.27±0.41 ^b	16.28±0.38 ^a	15.26±0.23 ^b
MCH	5.19±0.10	5.70±0.29	5.32±0.17	5.10±0.15	5.23±0.13	5.10±0.07
MCHC	33.43±0.35	33.14±0.26	33.15±0.19	33.47±0.27	32.65±0.37	33.53±0.51
WCC	16.68±1.07	20.03±2.10	16.96±1.32	14.50±1.28	16.29±2.38	17.55±1.31
Lymphocyte	10.87±0.98	13.54±1.56	13.74±1.07	11.64±1.10	10.36±2.04	10.32±0.80
Monocyte	0.53±0.10 ^{abc}	0.71±0.10 ^{ab}	0.27±0.06 ^c	0.28±0.08 ^{bc}	0.48±0.20 ^{bc}	0.93±0.29 ^a
Eosinophil	0.20±0.07	0.42±0.10	0.40±0.09	0.48±0.17	0.27±0.06	0.35±0.16
Platelet count	643.27±65.74 ^{ab}	1372.89±392.01 ^a	693.55±102.33 ^{ab}	140.0±53.11 ^b	503.50±192.01 ^b	554.25±65.03 ^b

^{a,b,c} Means with different superscripts on same row differ significantly ($p < 0.05$). SEM: Standard error of the means. WCC: white blood cell count; RBC: blood erythrocytes count; MCV: mean corpuscular volume; HCT: Haematocrit; MCH: mean corpuscular haemoglobin; MCHC: mean corpuscular haemoglobin concentration; Hb: haemoglobin. Day 0: initial values, Day 7: interval 1, Day 28: interval 2, Day 42 (VMO₀): interval 3 Vaccination+ 0% *M. oleifera* inclusion, Day 42 (VMO₂₀): interval 3 Vaccination+ 20% *M. oleifera* inclusion, Day 42 (VMO₅₀): interval 3 Vaccination+ 50% *M. oleifera* inclusion.

Table 4: Comparison of growth performance parameters across phases from day 0-42.

Variable	Adaptation phase	Vaccination phase	Vaccination+ <i>M. oleifera</i> phase (Day 29-42)		
	Day 1-6 (Mean±SEM)	DAY 7-28 (Mean±SEM)	VMO ₀ (Mean±SEM)	VMO ₂₀ (Mean±SEM)	VMO ₅₀ (Mean±SEM)
BW (Kg)	19.27±1.47 ^b	20.24±7.51 ^b	25.17±1.7 ^a	19.63±0.73 ^b	22.25±1.28 ^{ab}
BWG	0.55±0.25 ^{ab}	0.52±0.27 ^{ab}	0±0.26 ^b	1±0.27 ^{ab}	1.38±0.32 ^a
GR	0.08±0.04 ^{ab}	0.07±0.04 ^{ab}	0±0.04 ^b	0.14±0.04 ^{ab}	0.20±0.05 ^a
MWG	0.70±0.18 ^{ab}	1.13±0.13 ^a	0.33±0.21 ^b	0.92±0.23 ^{ab}	1.21±0.24 ^a
FI		7.13±2.05	7.42±3.57	6.57±4.88	6.59±3.93
FCR		0.04±0.00	0.03±0.00	0.03±0.00	0.03±0.00

^{a,b} Means with different superscripts on the same row differ significantly ($p < 0.05$). SEM: Standard error of means.

There were significant differences ($p < 0.05$) amongst growth performance parameters and no differences were observed ($p > 0.05$) for feed efficiency parameters (Table 4). VMO₅₀ displayed higher mean values on BWG, GR, and MWG ($p < 0.05$) and there were no significant differences ($p > 0.05$) in FI and FCR for VMO₀, VMO₂₀ and VMO₅₀ in the *M. oleifera* supplementation phase (Table 4). VMO₀ had 0 GR and BWG.

Haematological profiles may be utilised to assess the immunological status in goats (Al-Seaf and AlHarbi, 2012), however, factors such as age, nutrition, stress, management and environmental factors are known to have an effect on blood profiles in small ruminants (Mohammed et al., 2016). In comparison across the phases, there were no significant differences for total white blood cell counts (WCC) throughout the study. This varies from the findings of Jo et al. (2014), who observed an increase in total WCC following vaccination of indigenous goats. However, it agrees with Kumar et al. (2017), who found no significant differences in total WCC but observed significant differences in monocyte counts. Furthermore, in agreement with Kumar et al. (2017), monocyte counts significantly reduced following vaccination, this could have been due to a mild inflammation following vaccination. Useh et al. (2010), reported a significant decrease in monocyte count following blackleg infection, however, there was no significant difference in total WCC. Therefore, it may be useful to observe the specific leukocytes rather than WCC in separation (Cornell University, College of Veterinary Medicine, 2020). The results of the current study also revealed a significant increase of monocyte counts on VMO₅₀ compared to monocyte counts following the vaccination phase. The significant differences in monocyte counts observed in the literature and the present study show that monocytes act as first-line mediators that prepare the immune system for defence. Inclusion of *M. oleifera* commenced on day 22 post-vaccination to allow sufficient time for the animals to develop immunologic protection (Hoebe et al., 2004). Therefore, the significant increase in monocyte count for VMO₅₀ suggests an enhanced immune response to vaccination which could have been caused by 50% *M. oleifera* inclusion as compared to 0% and 20% inclusion levels. This observation could also be associated with the higher content of heavy metals such as Ag and Cd in the experimental diet due to supplementation with 50% *M. oleifera* as reported by Kar et al. (2015). However, Shen et al. (2019) stated that toxicity from heavy metals such as Cd affects the bones, thus, causing anaemia in the animals. However, there was no evidence of anaemia in animals that were fed a diet containing 50% *M. oleifera* although MCV and RBC levels were lower as compared to VMO₂₀.

Higher levels of platelet count outside the reference intervals as seen on day 0 are not of pathological importance

especially in young animals but sometimes maybe bought about by an infection or anaemia (Evstatiev et al., 2014). According to Kumar et al. (2017), platelet count does not play any role in immune response following vaccination. However, in the present study, mean platelet counts increased during the adaptation period and significantly reduced following vaccination towards the recommended reference ranges of healthy goats as recommended by Jackson and Cockcroft (2002). Although there were no significant differences in mean platelet counts during the *M. oleifera* supplementation phase amongst all treatments, 0% *M. oleifera* supplementation decreased platelets counts to below the recommended reference interval and supplementation with 20% and 50% *M. oleifera* leaves decreased the platelet counts to within the recommended reference interval of healthy goats. This could be due to the *M. oleifera* immune-boosting effects as reported Adouko et al. (2020). Metcalf-Pate et al. (2013) and Ali et al. (2015), revealed that platelets may play a role in the immune response in animals which agrees with the results of the present study. According to Brown et al. (2016b), low levels of Hb and RBC may also be an indication of anaemia, however, the results in this study showed values within the reference intervals. Therefore, we can safely rule out the possibility of anaemia in the present study and assume that supplementing with 20% and 50% of *M. oleifera* assisted in boosting the immune system of the animals.

Mean corpuscular volume (MCV) is a measurement of the average size of RBC. However, in the present study, MCV levels on day 0 were slightly below the reference range of healthy goats of (16–25 fL), this was also observed by Mohammed et al. (2016) in different goat breeds. The causes of these discrepancies are not clear, since the goats did not any show signs of anaemia. A significant increase to within the reference range of healthy goats in MCV for VMO₂₀ was expected due to higher levels of Cu and Fe minerals in the treatment diet which aid in red blood cell formation compared to treatment diets for VMO₀ and VMO₅₀ (Gaston et al., 2021). Therefore, it seems that anaemia may develop in the absence of these minerals. However, according to Wada (2004), high levels of Ca in the diet reduce the occurrence of anaemia even when Cu is deficient, therefore justifying the absence of anaemia in goats used in the present study.

Furthermore, the goats in VMO₂₀ were observed to be healthy, indicating that the higher Fe content than the recommended requirement for indigenous goats of 50–100mg Fe/kg did not cause any toxicity (Souza, 2014; Alfaro et al., 2021). The results of the present study are similar to the findings of Jiwuba et al. (2017), who observed significant differences in MCV values for *M. oleifera* inclusion levels and MCV levels were maintained within the range of healthy goats, suggesting that *M. oleifera*

may be used as a supplement without compromising the nutritional quality of the feed.

Balanced nutrition is vital in maintaining a healthy immune status in animals (Brown et al., 2016). Insignificant differences in most of the blood parameters among VMO₀, VMO₂₀, and VMO₅₀ observed in this study agree with some previous studies which reported insignificant differences in blood parameters of animals fed *M. oleifera* as a supplement (Moyo et al., 2012; Osman et al., 2012). Results of this study show that most of the blood parameters were within the range of healthy indigenous goats (Jackson and Cockcroft, 2002; Brown et al., 2016b; Daramola et al., 2005). However, VMO₀ had the lowest mean platelet and monocyte counts while VMO₅₀ had the highest counts. This could have been caused by a high content of minerals such as Ca in *M. oleifera* leaves which supports the formation of platelets, thus alleviating the occurrence of anaemia (Wada, 2004). According to Ali et al. (2015), there is a positive relationship between platelets and monocyte counts because platelets influence the immune response by activating immune cells such as monocytes. This is evident in the present study because as the level of *M. oleifera* inclusion increased, the means of platelet and monocyte counts increased as well. With this in mind, we can conclude that the presence of heavy metals and the low mineral content of Cu and Fe in the diets for VMO₅₀ in the present study did not bring about any toxicity or anaemia to the animals. Therefore, the possibility of anaemia in the present study can be ruled out and accept that supplementing with 20% to 50% of *M. oleifera* may assist in boosting the immune system of the animals.

Moringa oleifera is one of the alternative forages that are utilized for livestock nutrition in South Africa due to its high nutrient content (Qwele et al., 2013). Results from the present study showed that the growth performance parameters were influenced significantly by vaccination and varying *M. oleifera* inclusion levels. The mean values obtained for body weight gain, growth rate, and metabolic weight gain for VMO₅₀ were significantly higher than those in the control (VMO₀) but similar to VMO₂₀. The higher values for VMO₅₀ and VMO₂₀ on BWG, GR, and MWG could have been influenced by the lower levels of NDF and higher protein content in the diet, making it to be more digestible resulting in higher growth rates (Moyo et al., 2012). The significant decrease in BWG and GR following vaccination agrees with the findings from Jo et al. (2014), who reported a significant decrease in growth performance following vaccination of goats, which could be due to some of the side effects such as inflammation that may arise from vaccination. However, the introduction of varying levels of *M. oleifera* to the diets of the goats, increased BWG, GR, and MWG, suggesting

that *M. oleifera* due to its composition of elements such as antioxidants may aid in reducing inflammation and thus improving growth performance. This study agrees also with Mahfuz and Piao (2019); Aregheore (2002), who reported an improved growth performance in indigenous goats after supplementing with varying levels of *M. oleifera*. However, BWG and GR for VMO₀ were zero without affecting the FI, this agrees with the findings of Jo et al. (2014), who reported that vaccination decreases BWG without affecting FI. It could also be due to the significantly higher bodyweight of the goats in VMO₀ due to random grouping. The current findings disagree with the results of Yusuf et al. (2018), who reported insignificant differences in growth performance of indigenous goats fed 5 to 10% *M. oleifera* supplementation, therefore, it can be assumed that the minimum levels of *M. oleifera* that can be used as a supplement are 10% for improved growth performance in indigenous goats.

CONCLUSIONS AND RECOMMENDATIONS

In conclusion, supplementing with *M. oleifera* leaves in indigenous goat diets had a positive effect on growth performance and immunological response in goats following vaccination. Twenty percent and 50% *M. oleifera* supplementation yielded improved results as compared to the control diet following vaccination. Therefore, supplementation with 20% to 50% of *M. oleifera* may be used by farmers to overcome shortages of good quality feeds that may lead to malnutrition and poor immune responses in livestock. However, more research-based knowledge transfer is required for awareness to farmers especially in communal areas to produce *M. oleifera* in abundance for a more affordable protein and mineral source for their livestock.

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SCIENTIFIC CONTRIBUTION

The potential of *Moringa oleifera* as a feed supplement and an immunomodulatory in indigenous Pedi goats was discovered. The problem of poor immune responses in livestock following vaccination has been cited as one of the reasons for vaccine failure in communal livestock production systems. The study pursues an alternative feed source that may help to ease the challenges of communal livestock productivity after vaccination. Thereby, alleviating poverty levels and improving household food security.

The study explored the use of *Moringa oleifera* as a local resource to enhance the performance of the indigenous Ba-Pedi breed as compared to the majority of previous studies which evaluated *M. oleifera* in exotic breeds.

AUTHOR'S CONTRIBUTION

LG, KRM and TC conceptualization, visualization, funding acquisition, methodology, writing-review, and editing. LG and TC data collection and writing original draft preparation. LG, TLT and TC methodology investigation, formal analysis, visualization, writing-review, and editing. LG, KRM, TLT and TC writing-review, and editing, formal analysis, validation, resource.

DATA AVAILABILITY

The data that supports the findings of this study is available upon request.

CONFLICT OF INTEREST

The authors have declared no conflict of interest.

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