Carcass and meat traits, and non-carcass components of lambs fed ration containing increasing levels of urea

Características da carcaça, da carne e dos componentes não-carcaça de cordeiros alimentados com ração contendo níveis crescentes de ureia

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Abstract

The objective of this study was to evaluate the carcass and meat traits, and the non-carcass components of crossbred Dorper lambs fed diets with increasing levels of urea (0.0, 0.5, 1.0, and 1.5% of dry matter – DM). The experimental design was completely randomized with four treatments (urea inclusion levels) and six replicates per treatment. Lambs were fed *ad libitum* for 56 days and slaughtered at 37.9 ± 5.1 kg of body weight (BW). The weight and yield of carcass before and after cooling were not influenced by urea levels, with average values of 16.9 kg and 44.6% for cold carcass weight and yield. Urea levels did not affect the morphometric measurements, the fat deposition on the carcass, the weight of carcass cuts and the weight of non-carcass components. There was a quadratic effect of urea levels on the loin yield, which may achieve maximum value of 11.31% with the inclusion of 0.84% DM urea in the feed. The pH and the color coordinates L* (brightness), a* (red intensity) and C* (saturation) of the meat also showed quadratic response to the urea levels, where in the minimum value of 5.53 for pH, maximum value of 48.67 for L* and minimum values of 14.04 and 16.21 for a* and C* may be obtained by including 0.53 to 0.70% DM urea in the ration. The inclusion of 0.84% DM urea in the ration is recommended to obtain maximum yield of loin and meat with attractive characteristics to the consumer, which is characterized by high red intensity and brightness. If consumers have preference for lamb meat with a more intense red color, the inclusion of 1.5% DM urea should be considered in the ration formulation.
Key words: Color. Feedlot. Loin. Meat quality. Sheep.

Resumo

Objetivou-se com este estudo avaliar as características de carcaça, dos componentes não-carcaça e da carne de cordeiros mestiços Dorper alimentados com rações contendo níveis crescentes de ureia (0,0; 0,5; 1,0; 1,5% da matéria seca – MS). O delineamento foi inteiramente casualizado com quatro tratamentos (níveis de inclusão de ureia) e seis repetições por tratamento. Os cordeiros foram alimentados à vontade

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por 56 dias e abatidos com 37,9 ± 5,1 kg de peso corporal (PC). Os pesos e rendimentos de carcaça antes e após o resfriamento não foram influenciados pelos níveis de inclusão de ureia, com valores médios de 16,9 kg e 44,6% para peso e rendimento de carcaça fria. Os níveis de inclusão de ureia não afetaram as medidas morfométricas, a deposição de gordura e o peso dos cortes da carcaça; e o peso dos componentes não-carcaça. Houve efeito quadrático dos níveis de inclusão de ureia sobre o rendimento de lombo, que pode alcançar o valor máximo de 11,31% com a inclusão de 0,84% MS de ureia na ração. O pH e as coordenadas de cor L* (luminosidade), a* (intensidade de vermelho) e C* (saturação) da carne também apresentaram resposta quadrática aos níveis de inclusão de ureia, em que o valor mínimo de 5,53 para pH, valor máximo de 48,67 para L* e os valores mínimos de 14,04 e 16,21 para a* e C* podem ser obtidos com a inclusão de 0,53 a 0,70% MS ureia na ração. Recomenda-se a inclusão de 0,84% MS de ureia na ração para a obtenção de máximo rendimento de lombo e de carne com características qualitativas atrativas ao consumidor, que é caracterizada pela boa intensidade de cor vermelha e de brilho. Se os consumidores têm preferência por carne ovina com cor vermelha mais intensa, a inclusão de 1,5% MS de ureia deverá ser considerada na formulação da ração.

Palavras-chave: Confinamento. Cor. Lombo. Ovinos. Qualidade de carne.

Introduction

The consumption of lamb meat has increased considerably in recent years, mainly due to the supply of high quality meat obtained from young animals. In this aspect, the feedlot practice has been highlighted by the production of meat with superior quality (softness) when compared to that obtained from animals raised on pasture (TURINO et al., 2007), offer of standardized carcasses to the market, and high turnover and financial return (ALVES et al., 2014).

The production of high quality carcasses in feedlot depends on the high performance of lambs, which is maximized by providing properly balanced diets with energy and protein levels that favor high deposition of muscle tissue and enough fat deposition to meet consumer demand. On the other hand, this practice presents high production costs, once feeding is the most expensive component of the variable cost in animal production, representing between 80 and 90% (ALMEIDA et al., 2011). Among the ingredients used in large scale in ruminant nutrition, protein sources such as soybean meal and cottonseed meal are the most expensive (CLEMENTINO et al., 2007). Thus, the use of alternative protein sources is a way to minimize the costs of finishing lambs in feedlot.

Urea is an alternative food widely used in diets for ruminants (CANBOLAT; KARABULUT, 2010),

as a nitrogen source capable of supplying half the nitrogen requirements of rumen microorganisms, replacing conventional protein sources; due to its easy and rapid degradation in the rumen, promoting an efficient synthesis of microbial protein of high biological value; and low cost per unit of supplied protein (JENKINS et al., 2011). According to Chalupa (1968) for maximum urea utilization there must be 1% nitrogen in the total dry matter (DM) of diet, or 33% of the total dietary protein from non-protein nitrogen (NPN) of urea. However, the performance of feedlot lambs is not reduced when 1.7 to 1.95% of urea is added to the diet on DM basis (MAGALHÃES et al., 2006).

The carcass is the most valuable item in the commercialization of sheep. Different diets, its levels of energy and protein, and the finishing on pasture or in feedlot, among other factors, may affect the carcass traits (NÚÑEZ et al., 2007) and the development of non-carcass components (ANDRADE et al., 2009). Moreover, the inclusion of urea and different sources of true protein in the diet can influence the quality of the meat by modifying the chemical composition and growth of muscle tissues (ALVES et al. 2014).

Despite being described in literature the effects of feeding on a lambs carcass and meat characteristics, there is still little information about the use of alternative foods such as urea, especially

in intensive production systems. In this context, this study examined the influence that the inclusion of urea in the ration has on carcass and non-carcass traits, and meat quality of feedlot lambs.

Material and Methods

Experimental protocol

The research and procedures conducted on animals are in accordance with the Ethical Principles in Animal Experimentation adopted by the Colégio Brasileiro de Experimentação Animal (COBEA), and was approved by the Comissão de Ética no Uso de Animais (CEUA) of the Universidade Federal do Paraná (UFPR), Palotina Campus, with the protocol number 08/2012-CEUA issued by this institution.

The research was conducted at the Centro de Estudos em Pequenos Ruminantes (CEPER) of Palotina Campus, UFPR, located in Palotina, Paraná State, Brazil. Twenty four crossbred Dorper lambs were used with 25.0 ± 4.3 kg (average ± standard deviation – SD) of body weight (BW) and 3.5 months of age at the beginning of the experiment. The animals were identified with ear tags, weighed and housed in individual pens with 1.5 m² of area, slatted floor, automatic waterer and individual feeder.

The experimental design was completely randomized, with four treatments and six replications. The treatments were the inclusion of 0.0; 0.5; 1.0 and 1.5% urea in the ration on DM basis. The ration consisted of Tifton 85 hay (*Cynodon* spp) and concentrate feed (Table 1).

The adaptation period to the facility, management and diet was 15 days, and the trial period was 56 days under an environmental condition of 22.8 ºC (73.04 ºF) of temperature and 74.7% of relative humidity, on average. The rations were isoproteic with $17.03 \pm 0.13\%$ DM of crude protein (CP; Table 1).

Table 1. Proportion of ingredients and nutritional composition of the ration provided to feedlot lambs during the trial.

Composition [I]	Urea (% DM)			
	0.0	0.5	1.0	1.5
Tifton 85 hay (% DM)	36.0	40.0	34.0	25.0
Concentrate feed (% DM)	64.0	60.0	66.0	75.0
CPC (% DM) [II]	58.0	33.5	15.0	26.0
Soybean hulls (% DM)	5.0	25.0	49.0	19.0
Ground corn grain (% DM)	0.0	0.0	0.0	27.5
Mineral premix (% DM) [III]	1.0	1.0	1.0	1.0
Urea (% DM)	0.0	0.5	1.0	1.5
DM (%)	86.70	87.90	89.50	88.52
CP (% DM)	16.85	17.01	17.17	17.07
EE (% DM)	2.29	2.24	2.24	2.63
NDF (% DM)	28.55	44.75	56.63	32.71
Ca (% DM)	0.97	0.79	0.69	0.64
P (% DM)	0.46	0.35	0.27	0.36
Ca: P	2.10	2.26	2.56	1.88
TDN (% DM)	65.03	63.00	63.00	68.80
ME (Mcal kg DM^{-1})	2.36	2.27	2.27	2.48

[I] DM: dry matter; CP: crude protein; EE: ether extract; NDF: neutral detergent fibre, Ca: calcium; P: phosphorus; Ca: P: Ca: P ratio; TDN: total digestible nutrients; ME: metabolizable energy

[II] CPC – commercial pelleted concentrate: 87% DM; 18% CP; 2.5% EE; 9% of crude fibre; 10% of ash; 0.15% Ca; 0.06% P; 0.05% Na; 100 ppm BHT; 20 ppm Co; 45 ppm Cu; 55 ppm Fe; 10 ppm I; 50 ppm Mn; 0,3 ppm Se; 1,000 UI Vit A; 5,800 UI Vit D$_3$; 600 UI Vit E

[III] 1.2% Mg; 13.3% Na; 1% S; 6.5% P; 16.2% Ca; 2,250 ppm Mn; 86 ppm Cu; 1,400 ppm Fe; 200 ppm Co; 23 ppm Se; 4,500 ppm Zn; 177 ppm I; 100,000 UI Vit A; 65,000 Vit D$_3$; 60 UI Vit E.

Diets were fed as total mixed ration (TMR) and fractionated in two daily offers (08h00 and 14h00). Hay was shredded in particles of approximately 3 cm length to improve the utilization by the animals and reduce waste in the trough. The animals were fed *ad libitum* in the adaptation period and during the trial, keeping the orts in the trough at 10% of the amount of ration provided. Adjustments on the amount of ration provided were performed every five days based on the amount of orts.

On the first day of the trial period the animals were weighed after fasting (withdrawing feed for 12 h) in a scale with precision of 200 g. The trial period (56 days) was defined as the time required for the average animal slaughter BW to be reached, approximately 35 kg corresponding to the average BW for slaughtering lambs in the region (30 to 40 kg).

Assessments

The evaluation of the *in vivo* carcass composition was performed by measuring the ribeye area (REA_{US}) and backfat thickness (BFT_{US}) by ultrasound. The image capture was performed on the 30th and 56th day of the trial with the Landwind ultrasound model C40, equipped with electronic linear transducer (7 cm in length and frequency of 7.5 MHz). After shaving the wool and cleaning the skin in the region between the 12th and 13th thoracic vertebrae, the ultrasound images were captured using vegetable gel to provide better transducer coupling, conductivity and higher quality images. The transducer was equipped with an acoustic guide and was arranged perpendicularly to the length of *Longissimus lumborum* (*L. lumborum*), to enable the measurement of REA_{US} (cm²) and BFT_{US} (mm), the latter being measured at ¾ away from the medial position to the lateral position of muscle in relation to the midline. The images were identified with the ear tag number of each animal, digitalized with specific software that accompanies the equipment, and stored on a hard disk for further analysis and interpretation. The REA_{US} was calculated on the AutoCAD® software.

Upon reaching the preset slaughter weight, lambs were kept fasting (withdrawing feed for 16 h), weighed for recording the body weight at slaughter (SW) and sent to a commercial slaughterhouse in the region. The slaughter procedure was made by stunning the animals, followed by bleeding (performed by severing of the jugular veins and carotid arteries) and skinning to remove the skin. Blood was stored in a container and the head, paws and skin were removed and subsequently the four components were weighed individually. After, the gutting was performed followed by the collection of internal organs, which were sent to the Laboratório de Anatomia Patológica of Palotina Campus, UFPR. The internal organs were separated in heart, lungs + trachea, liver + gallbladder, spleen, kidneys, adipose deposits (omental, mesenteric, pelvic and perirenal fat) and gastrointestinal tract (GIT). The GIT was weighed with the digestive contents (DC) and later was individualized in rumen + reticulum, omasum, abomasum and intestine (small and large). All the organs were weighed individually, and after weighing the organs of GIT were emptied. The empty GIT organs were weighed and their yield relative to GIT and in relation to the empty body weight (EBW, obtained by the difference between SW and DC) was calculated.

After the slaughter, carcasses were weighed to record the hot carcass weight (HCW), identified and suspended by the metatarsal joints. The pH of the carcasses was measured 45 minutes after slaughter (pH_{PS}) with a pH meter introduced in *L. lumborum* muscle between the 4th and 5th lumbar vertebrae. Then the carcasses were transferred to a cold room at 4 °C, where they remained for 24 h. Post cooling, the pH was again measured (pH_{PC}) and the carcasses were weighed to record the cold carcass weight (CCW).

The SW, EBW, HCW and CCW were used to determine the biological yield [BY = (HCW/SW) x 100]; the hot carcass yield [HCY = (HCW/SW) x 100]; the commercial or cold carcass yield [CCY = (CCW/SW) x 100] and cooling loss [CL = (HCW – CCW)/HCW x 100].

The morphometry of the carcasses was evaluated based on the following measurements: carcass internal length (CIL) – measured with a measuring tape, corresponds to the distance from the front edge of the pubic bone to the cranial edge of the first rib; thoracic perimeter (TP) – using the bottom of the rib and the withers as basis, passing the measuring tape behind the shoulder; external leg length (ELL) – measured with a measuring tape in the lateral plane, corresponding to the distance between the base of the tail and the midpoint of the tarsal joint bones; internal leg length (ILL) – measured with measuring tape in the medial plane, corresponding to the distance between the front edge of the pubic bone and the midpoint of the tarsal joint bones; backfat thickness (BFT_{CARC}) taken on the external face of the *L. lumborum* muscle between the 12th and 13th thoracic vertebrae, using a digital caliper. The carcass compactness index (CCI) was calculated by the relation between CCW and CIL (kg cm^{-1}).

The carcasses were cut in the caudal-cranial direction into two halves, with the left half sectioned into seven commercial cuts as described by Colomer-Rocher and Espejo (1972): shoulder, leg, loin, uncovered ribs, true ribs, breast + flank and neck. In the dorsal portion of the *L. lumborum* muscle in the loin, by the cutoff point between the 13th thoracic vertebra and the 1st lumbar vertebra were measured the minimum and maximum thickness of fat on the ribeye (measure C and J, respectively). The transverse profile of the *L. lumborum* muscle was drawn on tracing paper to get the ribeye area (REA_{LOIN}), which was calculated by the ratio of the weight of the paper with a known area (4 cm²) and the weight of the paper where the ribeye profile was traced. The backfat thickness on the loin (BFT_{LOIN}) was obtained by averaging the measurements C and J.

Meat quality analyses were carried out in food analysis laboratories at the Pontifícia Universidade Católica do Paraná (PUC-PR), Campus São José dos Pinhais located in São José dos Pinhais, Paraná State. Loin samples were frozen at –18 °C for 90 days and later thawed to 5ºC for 24 h to determine the physical and chemical parameters of the meat which included pH, color, thawing loss (TL), water holding capacity (WHC) in raw samples; cooking loss (CKL) and shear force (SF) in cooked samples. Except for TL and WHC, these parameters were measured after thawing as described by Costa et al. (2011).

The pH measurement (pH_{MEAT}) was taken with a Jenway potentiometer (model 3020), calibrated to pH 4.0 and 7.0. The meat color was assessed on the surface of each sample with a colorimeter Minolta Chroma Meter (CR-200) calibrated to a standard white in the CIELab system, where the coordinates of brightness (L *), red coloration (a*), yellow coloration (b*), saturation (C*) and hue (H*) were measured. To determine the TL, the loin samples were thawed under refrigeration until reaching the internal temperature of 2 to 5°C. TL was calculated by the difference between the weight of samples before and after thawing, being expressed in percentage basis. The WHC was determined by approximately 2.0 g sample of each loin in triplicate following the methodology proposed by Hamm (1960). The samples were weighed and packed between two filter papers and acrylic plates, and received a pressure exerted by a weight of 10 kg for 5 min. After this process, the samples were weighed again and the amount of water lost was calculated. The result was expressed in percentage of water exuded on the weight of the initial sample.

To determine the CKL and SF, 24 loin samples were boiled in a water bath inside a plastic container resistant to heat until reaching an internal temperature of 70 °C and were subsequently cooled. The CKL corresponded to the weight difference between the samples before and after cooking, expressed as percentage basis. For the SF evaluation, three to seven portions were removed in a cylindrical shape of 1.27 cm diameter of each sample that was used to determine CKL. These portions were sheared perpendicularly to the direction of muscle fibers in the texturometer coupled to Warner-Bratzler device TA-XT Plus model, which measures the sample SF in kgf cm^{-2}.

Statistical analysis

Data were submitted to regression analysis (PROC REG) in which the level of inclusion of urea in the ration was considered the independent variable. Analyses were carried out to the second order, according to the model $\hat{Y}_{ij} = b_0 \text{ model}_0 + b_1 A i_1 + b_2 A i_2 + g_{(i,j)} + e_{(i,j)}$, where: $\hat{Y}_{ij}$ = value of dependent variable for j^{th} observation in the ith level of urea; b_0 = regression intercept; Ai = independent variable; b_1 = linear regression coefficient for the dependent variable; b_2 = quadratic regression coefficient for the dependent variable; $g_{(i,j)}$ = regression deviations for the jth observation in the ith level of urea; $e_{(i,j)}$) random error for the jth observation in the ith level of urea.

Pearson correlation analyses (PROC CORR) were performed to correlate the carcass traits among themselves and with the weights and yields of non-carcass components; and correlating the meat quality traits among themselves. In these analyzes, the *partial* option was used to adjust the correlation coefficients to the fixed effect of urea levels in the ration.

Statistical analyzes were performed using the *Statistical Analysis System*, version 9.0 (SAS, 2002). The level of 5% significance was adopted for all analyzes.

Results and Discussion

The levels of urea in the ration did not affect the (P> 0.05) quantitative traits of the lambs carcasses (Table 2). The average values for SW, EBW, HCW and CCW were 37.86; 33.21; 17.42 and 16.95 kg; and for BY, HCY, CCY and CL were 52.37; 45.92; 44.67 and 2.72%, respectively. Proper use of urea in ruminant feeding can promote the use of energy from carbohydrates and modify tissue deposition in carcasses (ALVES et al., 2014). Some studies have reported increased carcass weight when the protein level was raised, with linear increase until the inclusion of 1.4% of urea in the diet (MENDOZA JÚNIOR et al., 2007). However, as observed in this study (Table 2), some studies have shown that increasing the levels of urea in the diet do not affect the carcass traits of lambs (SOUZA et al., 2004b; VOLTOLINI et al., 2010). It is likely that these results are due to similar conditions for the development of the carcasses, in which the SW of the animals was similar in all diets assessed; and urea inclusion levels in diets being too low to cause any significant tissue changes in the carcasses.

The results for EBW, HCW and CCW in relation to the urea levels are in agreement with previous studies, which evaluated the levels of urea in the diet up to 1.2% (SOUZA et al., 2004b) and 1.95% (MAGALHÃES et al., 2006), and the replacing of conventional urea by slow-release urea (ALVES et al., 2014). On the other hand, Canbolat and Karabulut (2010) observed increased CCW with increased energy, protein and urea levels (0, 6, 12 and 18 g animal^{-1} d^{-1}).

Table 2. Means and coefficients of variation (CV) for carcass traits of feedlot lambs fed rations with increasing levels of urea.

Variable [I]	Urea (% DM)				CV	P-value [II]	
	0.0	0.5	1.0	1.5	(%)	L	Q
SW (kg)	37.67	38.08	38.00	37.73	13.40	0.9869	0.8813
EBW (kg)	32.94	32.78	33.38	33.68	13.97	0.7532	0.9165
HCW (kg)	17.43	17.05	17.56	17.59	15.49	0.8515	0.8750
CCW (kg)	16.93	16.67	17.13	17.01	15.41	0.8936	0.9665
BY (%)	52.87	51.78	52.54	52.19	3.44	0.6852	0.6749
HCY (%)	46.17	44.51	46.15	46.61	4.76	0.5083	0.2902
CCY (%)	44.83	43.54	45.03	45.09	4.76	0.6012	0.5072
CL (%)	2.94	2.18	2.44	3.22	43.27	0.6531	0.1292

[I] SW: slaughter weight; EBW: empty body weight; HCW: hot carcass weight; CCW: cold carcass weight; BY: biological yield; HCY: hot carcass yield; CCY: cold carcass yield; CL: cooling losses
[II] L: linear regression; Q: quadratic regression.

The average value for BY was similar to that described by Souza et al. (2004b) (average 51.0%) with addition of up to 1.2% of urea in the diet. Values lower than that obtained in this study were described by Ziguer et al. (2012) (average 47.2%), which used soybean hulls associated with different sources of non-protein nitrogen (conventional urea, protected urea, protected urea + urea conventional).

Additional results of this study obtained by Vivian et al. (2017) indicated there was no significant effect (P> 0.05) of urea levels on DM intake (DMI). The average DMI during the trial was 1.174 kg animal^{-1} d^{-1}, and met the requirements for fast and moderate growth according to the NRC (1985) (DMI = 1.0 to 1.3 kg DM animal^{-1} d^{-1}). Thus, the urea levels in the diet did not affect the weight of DC and did not cause depletion of food intake by physical filling of the GIT, which are factors that affect directly the BY.

The mean values for HCY and CCY are in agreement with the literature (46.0% and 44.5% respectively). The CL in sheep carcasses may vary between 1 and 7%, usually found close to 2.5%. This characteristic is influenced by the, sex, weight, fat covering of the carcass, temperature and humidity in the cold storage chamber, and the handling of the carcasses (MORENO et al., 2008). The muscle fiber composition is also of particular importance because muscles with predominance of red fibers are more susceptible to shortening by the cooling. This occurs due to red fibers have small ability to retain calcium and pH at low temperatures (ZEOLA et al., 2007). However, the muscle fibers are formed during pregnancy and only minor muscle fibers suffer from environmental interference, such as maternal nutrition (ZÜNDT et al., 2006). The CL values obtained in this study are close to the normal value observed in lamb carcasses, and are justified by similar deposition of fat in the carcass and its proper handling and storage immediately after slaughter.

The levels of urea did not affect (P> 0.05) the morphometric traits (Table 3), with average values of 67.4 cm, 72.3 cm, 45.1 cm; 36.3 cm and 0.255 kg cm^{-1} for CIL, TP, ELL, ILL and CCI, respectively. These traits are in agreement with previous studies that reported that the carcass morphometry is not influenced by its food system, as long as the animals are slaughtered in the same weight range (ALVES et al., 2014). The average CCI obtained in this study (0.255 kg cm^{-1}) was similar to that described by Souza et al. (2004b), who used up to 1.2% of urea in the diet of the lambs.

Table 3. Means and coefficient of variation (CV) for the morphometric measurements of the carcasses of feedlot lambs fed rations with increasing levels of urea.

Variable [1]	Urea (% DM)				CV	*P*-value [II]	
	0.0	0.5	1.0	1.5	(%)	L	Q
CIL (cm)	66.2	69.0	67.3	67.7	3.68	0.4672	0.3411
TP (cm)	70.8	72.3	73.3	72.9	5.05	0.2877	0.5502
ELL (cm)	44.5	45.4	44.7	45.8	6.52	0.5467	0.8879
ILL (cm)	35.3	36.9	36.1	37.1	5.82	0.2517	0.7882
CCI (kg cm^{-1})	0.255	0.261	0.255	0.251	12.32	0.7773	0.7684

[1] CIL: carcass internal length; TP: thoracic perimeter; ELL: external leg length; ILL: internal leg length; CCI: carcass compactness index
[II] L: linear regression; Q: quadratic regression.

The backfat thickness and ribeye area that were measured *in vivo* by ultrasound, in the carcass right after the slaughter and in the *L. lumborum* muscle were not affected (P> 0.05) by urea levels in the ration (Table 4). It was recorded mean values of 8.99 and 10.40 cm² for initial and final REA_{US}; 3.63 and 5.30 mm for initial and final BFT_{US}; 2.52 mm for BFT_{CARC}; 14.26 cm² and 3.61 mm for REA_{LOIN} and BFT_{LOIN}, respectively.

Table 4. Means and coefficient of variation (CV) for ribeye area (REA) and backfat thickness (BFT) measured *in vivo* by ultrasonography, directly in the carcass and in the *Longissimus lumborum* muscle of feedlot lambs fed ration with increasing levels of urea.

Variable [I]	Urea (% DM)				CV	*P*-value [II]	
	0.0	0.5	1.0	1.5	(%)	L	Q
Initial REA_{US} (cm²)	9.79	8.92	8.22	9.01	18.04	0.3141	0.2181
Final REA_{US} (cm²)	11.04	10.51	10.20	10.08	13.65	0.3064	0.7738
Initial BFT_{US} (mm)	3.87	3.54	3.48	3.60	14.57	0.3768	0.3320
Final BFT_{US} (mm)	5.72	4.72	5.56	5.17	17.83	0.5915	0.4782
BFT_{CARC} (mm)	2.44	2.26	2.82	2.56	28.51	0.5391	0.8842
REA_{LOIN} (cm²)	14.60	13.25	14.09	14.95	16.53	0.6940	0.2945
BFT_{LOIN} (mm)	3.64	3.43	3.59	3.77	25.02	0.7381	0.6382

[I] REA_{US}: ribeye area measured by ultrasonography; BFT_{US}: backfat thickness measured by ultrasonography; BFT_{CARC}: backfat thickness measured in the carcass; REA_{LOIN}: ribeye area measured in the *Longissimus lumborum* muscle; BFT_{LOIN}: backfat thickness measured in the *Longissimus lumborum* muscle
[II] L: linear regression; Q: quadratic regression.

Fat is the largest carcass component variation, increasing with age and weight at slaughter (MACEDO et al., 2000). This can be seen in the values obtained by ultrasound, in which the fat thickness on the REA increased during the trial regardless of urea levels in the ration (mean for initial BFT_{US} = 3.63 mm; mean for final BFT_{US} = 5.30 mm; Table 4). REA was not influenced by urea levels in the ration, but increased during the trial as was indicated by measurements made by ultrasound (mean for initial REA_{US} = 8.99 cm², mean for final REA_{US} = 10.40 cm²; Table 4).

The carcass evaluation made by ultrasound pre-slaughter resulted in higher values of fat thickness and a smaller REA compared to those done in the carcass (mean for BFT_{CARC} = 2.52 mm) and loin (mean of 3.61 mm for BFT_{LOIN}; mean of 14.26 cm² for REA_{LOIN}; Table 4). This was also reported by Ítavo et al. (2009), which suggested that the existence of wool, the softness of the fat and the mobility of the skin may be the reasons for the lower accuracy of the measurements performed by ultrasound. Nevertheless, the final REA_{US} showed moderate and positive correlation with REA_{LOIN} (r = 0.69; P = 0.0008) and BFT_{LOIN} (r = 0.50; P = 0.0245), while the final BFT_{US} showed no correlation (P> 0.05) with the measurements in the carcass and in the loin. Thus, even with low accuracy, the REA measured by ultrasound in pre-slaughter can be an indicator of muscle development and fattening degree of lamb carcasses.

A minimum fat cover is needed to protect the carcass from water loss and cold burns during cooling and freezing, and to keep the organoleptic characteristics of the meat (SANTOS et al., 2009). The average fat thickness covering the carcasses which was obtained in this study (BFT_{CARC} = 2.52 mm) was higher than that reported by Ítavo et al. (2009). The mean value for REA measured in the loin (REA_{LOIN} = 14.26 cm²) was close to that reported by Urano et al. (2006), who observed a REA of 14.8 cm² in lambs slaughtered at 37.7 kg BW. These

results indicate that the SW and the feedlot period were enough to produce standardized and well-developed carcasses and, with no influence of urea levels in the diet on the deposition of covering fat.

The weight of the commercial cuts were not affected (P> 0.05) by urea levels in the ration (Table 5). The mean values for weight of neck, uncovered ribs, true ribs, breast + flank, loin, shoulder and leg were 0.719; 0.503; 0.949; 1.110; 0.924; 1.562 and 2.784 kg, respectively.

For the yield of cuts, there was a quadratic effect of urea levels (P <0.05) on the loin yield, which was higher in lambs that received ration with 1% DM urea (Table 5). This can be explained by the increase of CP in the diet promoted by this level of inclusion of urea. The regression equation indicates that the maximum loin yield (11.31%) can be obtained by including 0.84% DM urea in the ration. This is interesting from an economic point of view, since it is possible to increase the yield of a highly valuable commercial prime cut by supplying a cheaper ration for the lambs. Other studies showed no effect of urea levels in the diet on the loin yield, and on the weight and yield of other commercial cuts of the lamb carcass (SOUZA et al., 2004b; VOLTOLINI et al., 2010).

The mean values of yield of neck, uncovered ribs, true ribs, breast + flank, loin, shoulder and leg were 8.35; 5.83; 11.05; 12.95; 10.84; 18.30 and 32.68%, respectively. It is noteworthy that the yield of loin, shoulder and leg, which are the prime cuts of higher commercial value, accounted for 61.83% of the lambs carcass. From these cuts, leg showed greater weight (mean of 2.784 kg) and higher yield (mean of 32.70%), which confirms that the leg is the noblest cut of the carcass because of the greater amount of muscle mass and, consequently, the higher meat yield (RIBEIRO et al., 2009).

Table 5. Means and coefficient of variation (CV) for weight and yield of carcass cuts of feedlot lambs fed rations with increasing levels of urea.

Variable [I]	Urea (% DM)				CV	P-value [II]	
	0.0	0.5	1.0	1.5	(%)	L	Q
Weight (kg)							
Neck	0.718	0.682	0.710	0.760	26.74	0.6846	0.6150
Ribs$_{UNCOV}$	0.518	0.482	0.485	0.522	24.90	0.9608	0.5090
Ribs$_{TRUE}$	0.941	0.924	0.974	0.951	19.74	0.8272	0.9543
BF	1.138	1.062	1.071	1.163	18.61	0.8472	0.3506
Loin	0.884	0.931	0.958	0.925	14.67	0.5497	0.4947
Shoulder	1.567	1.540	1.568	1.571	15.91	0.9360	0.8970
Leg	2.849	2.776	2.753	2.756	14.76	0.6950	0.8340
Yield (%)							
Neck	8.31	8.08	8.30	8.66	16.28	0.6302	0.6306
Ribs$_{UNCOV}$	5.92	5.73	5.65	6.00	14.39	0.9269	0.4562
Ribs$_{TRUE}$	10.86	10.95	11.41	10.95	8.02	0.6655	0.4413
BF	13.22	12.58	12.55	13.41	7.36	0.7827	0.0623
Loin [III]	10.31	11.18	11.24	10.70	6.71	0.3396	0.0167
Shoulder	18.19	18.39	18.44	18.20	5.50	0.9537	0.6142
Leg	33.21	33.10	32.41	32.09	6.39	0.3054	0.9206

[I] Ribs$_{UNCOV}$: uncovered ribs; Ribs$_{TRUE}$: true ribs; BF: breast + flank
[II] L: linear regression; Q: quadratic regression
[III] $Y_{LOIN} = -1.4042U^2 + 2.3547U + 10.3186$ (R^2 = 0.9972).

There was no significant effect (P> 0.05) of urea levels on the weight of head, paws, skin, red viscera and fatty deposits (Table 6). The mean values for weight of head, paws, skin, blood and adipose deposits were 2.177; 0.918; 4.371; 1.934 and 0.929 kg, respectively. The mean values for weight of heart, lungs, liver, spleen and kidneys, which are characterized as red viscera, were 0.160; 0.660; 0.859; 0.171 and 0.132 kg, respectively. When the organ weights are uniform and have an earlier maturation, they are less affected by dietary treatment in a later stage of life (FASAE et al., 2011). Heart (mean of 0.160 kg) and lungs (mean of 0.660 kg) weights are in agreement with other studies (MORENO et al., 2011), demonstrating that these organs maintain the integrity and have priority in the use of nutrients, regardless of their diet. The lack of effect of urea levels on adipose deposits was also reported in other studies evaluating the weight of non-carcass component in lambs receiving up to 1.4% urea in the diet (MEDEIROS et al., 2011).

High weight of some non-carcass components such as head, skin and blood can adversely affect the carcass yield (LANDIM et al., 2007), which did not occur in this study. In fact, the head, paws and skin weights, the assembly formed by these three components (HPS) and the blood weight showed no correlation (P> 0.05) with BY, HCY and CCY, which confirms that these components did not affected the carcass yield of the lambs.

Except for the yield of rumen + reticulum in relation to the empty body weight (RR_{EBW}), there was no effect (P> 0.05) of the urea levels in the ration on weight and yield of GIT and its organs, and also on the DC (Table 7).

Table 6. Means and coefficient of variation (CV) for weight of head, paws, skin, blood, red viscera and fatty deposits of feedlot lambs fed rations with increasing levels of urea.

Variable [I]	Urea (% DM)				CV (%)	P-value [II]	
	0.0	0.5	1.0	1.5		L	Q
Head (kg)	1.973	2.219	2.223	2.267	13.61	0.1290	0.4449
Paws (kg)	0.871	0.904	0.925	0.968	14.48	0.2047	0.9279
Skin (kg)	4.016	4.785	4.167	4.584	15.45	0.3591	0.6266
HPS (kg)	7.021	7.908	7.315	7.819	11.54	0.2967	0.7052
Blood (kg)	1.833	1.768	2.174	1.935	32.74	0.5642	0.7125
Heart (kg)	0.157	0.158	0.165	0.159	14.25	0.7726	0.7218
Lungs (kg)	0.626	0.634	0.689	0.687	17.85	0.2839	0.8995
Liver (kg)	0.891	0.843	0.792	0.909	18.64	0.9944	0.2218
Spleen (kg)	0.184	0.162	0.171	0.167	27.64	0.6267	0.6920
Kidneys (kg)	0.128	0.135	0.139	0.125	15.52	0.9047	0.2570
Red viscera (kg)	1.985	1.931	1.955	2.047	14.59	0.7114	0.5646
Fatty deposits (kg)	0.812	0.850	1.040	0.981	42.16	0.3728	0.7511

[I] HPS: sum of head, paws and skin weights; Red Viscera: sum of heart, lungs, liver, spleen and kidneys weights; Adipose Deposits: sum of omental, mesenteric and perirenal fat weights
[II] L: linear regression; Q: quadratic regression.

Table 7. Means and coefficients of variation (CV) for weights of gastrointestinal tract (GIT), digestive content (DC) and organs from GIT and their yields relative to GIT and to the empty body weight (EBW) of feedlot lambs fed rations with increasing levels of urea.

Variable [I]	Urea (% DM)				CV	*P*-value [II]	
	0.0	0.5	1.0	1.5	(%)	L	Q
Full GIT (kg)	7.257	8.138	7.428	6.840	14.73	0.3657	0.1212
DC (kg)	4.724	5.297	4.622	4.053	20.54	0.1403	0.1582
Empty GIT (kg)	2.533	2.840	2.806	2.787	14.75	0.3201	0.3591
RR (kg)	0.763	0.884	0.874	0.847	12.56	0.2019	0.0935
Omasum (kg)	0.095	0.120	0.096	0.100	21.74	0.9302	0.3407
Abomasum (kg)	0.234	0.219	0.225	0.228	19.57	0.8721	0.6484
Intestine (kg)	1.441	1.618	1.611	1.612	20.38	0.3878	0.5373
RR_{GIT} (%)	30.23	31.37	31.51	30.71	10.58	0.7882	0.4991
$Omasum_{GIT}$ (%)	3.73	4.22	3.45	3.60	16.71	0.3579	0.6083
$Abomasum_{GIT}$ (%)	9.29	7.83	8.10	8.18	18.98	0.2841	0.2701
$Intestine_{GIT}$ (%)	56.75	56.58	56.94	57.52	7.42	0.7441	0.8434
RR_{EBW} (%) [III]	2.35	2.71	2.64	2.53	11.32	0.3606	0.0483
$Omasum_{EBW}$ (%)	0.29	0.37	0.29	0.30	19.50	0.6385	0.2009
$Abomasum_{EBW}$ (%)	0.72	0.67	0.69	0.68	21.19	0.6710	0.7110
$Intestine_{EBW}$ (%)	4.44	4.97	4.84	4.77	16.97	0.5476	0.4027
GIT_{EBW} (%)	7.80	8.72	8.46	8.27	11.77	0.5007	0.1980

[I] GIT: gastrointestinal tract; DC: digestive content; RR: rumen + reticulum; EBW: empty body weight
[II] L: linear regression; Q: quadratic regression
[III] $RR_{EBW} = -0.4713U^2 + 0.8020U + 2.3645$ ($R^2 = 0.9009$).

There was a quadratic effect (P <0.05) of urea levels on RR_{EBW}, which was higher (2.71%) in lambs that received ration with 0.5% DM of urea (Table 7). Under the conditions of this study, the regression equation indicated that the inclusion of 0.85% DM of urea in the ration allows achieving the maximum value of RR_{EBW}, which corresponds to 2.71% and is similar to the value found for the inclusion of 0.5% DM of urea in the ration. Thus, the maximum yield of rumen + reticulum can be achieved by providing diets containing 0.5 to 0.85% DM of urea to the lambs.

The increase in the forage:concentrate ratio (F:C) in the diet, which leads to increased fiber content and lower energy content in its composition, leads to increased size of the rumen, reticulum and omasum (REIS; SILVA, 2006). Although the rations were isoproteic (17.03% DM of CP on average), the F:C and the neutral detergent fiber (NDF) content varied among the experimental diets (Table 1). The F:C ratio ranged from 25:75 to 40:60 and the NDF content ranged from 28.55 to 56.63%, where the rations with 0.5 and 1.0% DM of urea showed higher NDF (44.75 and 56.63%, respectively). The high NDF on these diets is associated with the greater inclusion of soybean hulls in their compositions with inclusion of 25 and 49% DM, respectively. In assessing the nutrient intake by lambs of this study, Vivian et al. (2017) observed a quadratic effect of urea levels on NDF intake, which reached a maximum value (0.665 kg DM d⁻¹) in the ration with 1.0% DM of urea. However, the regression equation indicates that the maximum NDF intake (0.604 kg DM d⁻¹) may be achieved with the inclusion of 0.85% DM of urea in the ration. This justifies the possibility of obtaining the maximum RR_{EBW} (2.71%) when the ration containing the same inclusion of urea is provided to the lambs. These results show that the type of food and the F:C ratio of the diet has great influence on the non-carcass components, especially those involved in the digestion and absorption of nutrients.

Although they were not influenced (P> 0.05) by urea levels, the pH_{PS} values were higher in the carcasses of lambs that received rations without urea (6.68) and with 0.5% DM of urea (6.83); and pH_{PC} value was higher in the carcasses of lambs fed diets with 0.5% DM of urea (6.05; Table 8). The normal values for sheep meat range from 7.3 to 7.5 at slaughter, resulting in a final value of 5.5 to 5.8 between 12 to 24 h after slaughter (ZEOLA et al., 2007).

The carcass pH can be influenced by the SW and the BFT (CEZAR; SOUSA, 2007). The pH_{PS} and pH_{PC} showed no correlation (P> 0.05) with the SW and the BFT measured by ultrasound in the pre-slaughter (final BFT_{US}), in the carcass (BFT_{CARC}) and loin (BFT_{LOIN}), which indicates that these traits did not affect the carcass pH of the lambs. The greater supply of protein in the diet, which affects muscle chemical composition, may determinate high pH in the carcass (OSÓRIO et al., 2009). The diets used in this study were isoproteic, however, it was expected that the increasing levels of urea could change the pH_{PS} and pH_{PC}, which was not proved (Table 8).

Although the pH_{MEAT} is within the normal range (5.5 to 5.8), it showed a quadratic response to urea levels in the ration (Table 8). The lowest pH values were recorded in the meat of lambs that received rations with 0.5 and 1.0% DM of urea (5.55), and the highest in the lambs fed ration with 1.5% DM of urea (5.68). The regression equation indicates that the minimum value for pH_{MEAT} (5.53) can be obtained by including 0.53% DM of urea in the diet.

Gluconeogenesis and ureogenesis are vital for the metabolism of ruminants. However, ammonia affects the gluconeogenesis, reducing the conversion of propionate to glucose. This mechanism is still not well described, but significant effects have been detected suggesting a specific response of ammonia in the competition for energy (ATP) required for synthesis of glucose or urea (NORO et al., 2012). The small glycogen reserve causes the decrease of glucose levels in the muscles, which determines lower production of lactic acid by anaerobiosis and hence little decrease in pH (ZEOLA et al., 2007). Probably the darkest meat from lambs fed ration with 1.5% DM of urea can be associated with the highest concentration of ammonia in the muscle, which may have interfered in gluconeogenesis, reducing the availability of glucose to the formation of glycogen reserves in the muscles. This determined the highest pH in the meat (pH_{MEAT} = 5.68) of lambs receiving 1.5% of DM urea in the ration (Table 8).

The pH is directly related to other meat quality traits, which was confirmed by the moderate to high correlations of pH_{MEAT} with coordinates L*, a* and C*. The sheep meat usually has values from 30.03 to 49.47 for L*; 8.24 to 23.53 for a* and from 3.38 to 11.10 for b* (SOUZA et al., 2004a). Thus, the values found in this study (Table 8) are close to those described in the literature.

There was a quadratic effect (P <0.05) of urea levels on the color coordinates L*, a* and C* (Table 8). The b* and H* coordinates were not (P> 0.05) influenced by urea levels and showed mean values of 8.30 and 28.71, respectively.

Table 8. Means and coefficients of variation (CV) for carcass pH and meat quality traits of feedlot lambs fed rations with increasing levels of urea.

Variable [I]	Urea (% DM)				CV (%)	P-value [II]	
	0.0	0.5	1.0	1.5		L	Q
pH_{PS}	6.68	6.83	6.46	6.56	4.31	0.1854	0.8975
pH_{PC}	5.80	6.05	5.86	5.83	5.75	0.9274	0.3729
TL (%)	2.93	2.18	2.08	2.30	63.48	0.4749	0.4629
WHC (%)	34.16	35.24	34.21	34.50	8.68	0.9867	0.7921
pH_{MEAT} [III]	5.57	5.56	5.55	5.68	1.56	0.0485	0.0291
CKL (%)	25.27	24.98	23.75	24.22	13.91	0.5003	0.7823
SF (kgf cm^{-2})	2.81	2.86	3.01	2.58	21.52	0.6450	0.3537
Colour coordinates							
L* [III]	46.50	48.96	47.03	42.53	8.81	0.0689	0.0269
a* [III]	15.48	14.43	14.17	16.52	10.07	0.3375	0.0040
b*	8.77	8.37	7.87	8.20	14.09	0.3148	0.4465
C* [III]	17.84	16.70	16.23	18.52	9.18	0.6284	0.0065
H*	29.44	30.13	28.94	26.58	12.22	0.1378	0.2962

[I] pH_{PS}: pH of carcass post-slaughter; pH_{PC}: pH of carcass post-cooling; TL: thawing loss; WHC: water holding capacity; pH_{MEAT}: pH of the meat; CKL: cooking loss; SF: shear force; L*: brightness; a*: intensity of red colour; b*: intensity of yellow colour; C*: colour saturation; H*: colour tone of the meat
[II] L: linear regression; Q: quadratic regression
[III] $pH_{MEAT} = 0.1442U^2 - 0.1529U + 5.5749$ ($R^2 = 0.9405$)
  $L* = -6.9145U^2 + 7.6135U + 46.5787$ ($R^2 = 0.9924$)
  $a* = 3.4414U^2 - 4.5812U + 15.5618$ ($R^2 = 0.9529$)
  $C* = 3.4862U^2 - 4.9067U + 17.9347$ ($R^2 = 0.9313$).

The inclusion of 0.5% DM of urea in the ration determined a clearer meat by having greater L* value (48.96), while adding 1.5% DM of urea resulted in darker meat, with lower L* (42.53; Table 8). Although higher brightness was observed in the meat of lambs fed ration with 0.5% DM of urea, the regression equation indicates that the maximum value of 48.67 in L* can be obtained with the inclusion of 0.55% DM of urea in the ration.

The lowest and the highest value for a* were recorded in the diets with 1.0 and 1.5% DM of urea (14.17 and 16.52; Table 8), which characterized lower and higher intensity of red color in the meat of lambs that received these rations, respectively. Based on the regression equation, the minimum value of 14.04 to a* can be obtained by including 0.67% DM of urea in the ration.

The coordinate C* showed a similar response pattern to the coordinate a* relative to the urea levels, with lower and higher value for 1.0 and 1.5%

DM of urea inclusion (16.23 and 18.52 respectively; Table 8). Thus, the meat had become brighter and more matte in lambs receiving diets with 1.0 and 1.5% DM of urea, respectively. The value of C* for the inclusion of 1.0% DM of urea was very close to the minimum value estimated for this trait using the regression equation, which was 16.21 with the inclusion of 0.70% DM of urea in the ration.

The meat of the lambs used in this study was redder and brighter, with higher values for the coordinate a* and lower values for coordinate C* when compared to results obtained by Souza et al. (2004a). Some studies suggest that the type of food can change the color of the meat. The greater intake of forage by grazing animals stimulates the increase of myoglobin levels in the muscles due to its higher levels of carotenoids (MORENO et al., 2008). Otherwise, some reports comment that the nature of the food (forage or grain) has little influence due to intense transformations that food suffers in the rumen (OSÓRIO et al., 2009). In this

study, the darker (lower L*), redder (higher a*) and more matte (higher C*) profile of meat from lambs fed ration with 1.5% DM of urea was determined by pH, as described above. Probably these lambs had few glycogen stores in their muscles, which do not allow the meat to reach a pH low enough to determine normal color, regardless of animals age and meat tenderness (ZEOLA et al., 2007).

The measurement of meat color by the objective method determined that greater values for L* are associated with paler meat, while greater values for a* and b* indicate a redder and yellower meat, respectively. The L* coordinate is associated with the amount of water in the tissue and the progress of post mortem biochemical reactions. The coordinate a* is an indicator of the color intensity and is related to the muscle oxymyoglobin content. The color of meat measured in relative values to the light reflection, represented by L*, is inversely proportional to the concentration of myoglobin in the muscle tissue, thus, the higher concentration of myoglobin reflects the smaller L* value (ODA et al., 2004). This must have occurred in the muscle tissue of lambs fed ration with 1.5% DM urea, as they produced meat with greater a* value and lower value of L*.

In general, lambs fed ration with 1.5% DM of urea showed a darker (lower L*), redder (higher a*) and more matte (higher C*) meat. These characteristics may be associated with pH_{MEAT} that regardless of being within the normal range was higher in meat from lambs fed ration with 1.5% DM of urea. The correlation analysis indicated that the pH_{MEAT} showed moderate and positive correlation with the coordinates a* (r = 0.44; P = 0.0472), thus, the increase of pH_{MEAT} leads to a more intense red color. It was also found that the coordinate a* showed moderate and negative correlation with the L* coordinate (r = –0.56; P = 0.0082), and high and positive correlation with C* coordinate (r = 0, 95; P <0.0001). Therefore, the increased intensity of red color causes the decrease of brightness and increase of saturation, resulting in darker and more matte meat.

The TL, WHC, CKL and SF were not affected (P> 0.05) by urea levels in the ration (Table 8), and showed mean values of 2.38%, 34.50%, 24.53% and 2.81 kgf cm^{-2}, respectively. The pH can also affect other meat quality traits, such as TL, WHC, CKL and SF. Fernandes Júnior et al. (2013) observed higher CKL (9.58%) in the meat of animals fed diets with high content of sunflower cake (33.8% DM of total diet), due to the lower pH of the meat (5.37). Except for CKL, the pH_{MEAT} showed no correlation (P> 0.05) with the meat quality traits. There was a moderate and positive correlation between pH_{MEAT} and CKL (r = 0.43; P = 0.0494), indicating that the increase of pH_{MEAT} causes an increase in CKL, therefore less water bound to myoglobin, which contradicts the results of Fernandes Júnior et al. (2013). As the CKL is not affected by urea levels in the ration (Table 8), it is likely that the correlation between pH_{MEAT} and CKL are associated with some individual variation in animals or meat samples that were analyzed.

The mean value for SF indicated that the meat of the lambs showed intermediate softness, because sheep meat with SF lower than 2.27 kgf cm^{-2} are classified as soft, and between 2.28 and 3.63 kgf cm^{-2} are classified as intermediately soft (CEZAR; SOUSA, 2007). Given the lack of effect of urea levels on the SF (Table 8), the results for this trait were expected, since the animals were from similar genetic groups and slaughtered with the same SW. Whereas SF indicates the tenderness of the meat, it showed intermediate softness (mean of 2.82 kgf cm^{-2}), with adequate quality for commercialization.

Conclusion

The inclusion of up to 1.5% of urea in the ration, based on dry matter (DM), does not increase the carcass weight and yield, and the carcass compactness; and the weight of commercial cuts and non-carcass components of feedlot lambs.

The inclusion of 0.84% DM of urea in rations for lambs is recommended to obtain maximum

loin yield and meat with characteristics that are attractive to the consumer, which is characterized by the good intensity of red color and brightness. However, if consumers have a preference for lamb with a more intense red color, the inclusion of 1.5% DM of urea should be considered in the formulation of the ration.

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