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# Impact of water restriction periods on carcass traits and meat quality of feedlot lambs in the Brazilian semi-arid region

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#### ABSTRACT

Water restriction periods were evaluated in crossbred lambs (n = 32) distributed in one of four treatments: without water restriction, water restriction for 24, 48 and 72 h. The water restriction for 72 h reduced the water and dry matter intakes, body weight at slaughter and hot and cold carcass yields. Water restriction did not affect the weight of the carcass cuts and the chemical composition of the meat. The fatty acid EPA increased and DHA reduced with increasing water restriction period. There was an increasing linear effect for meat shear force, with less force (30.5 N/cm<sup>2</sup>) for sheep meat without water restriction and higher force (45.8 N/cm<sup>2</sup>) for those with water restriction for 72 h. The period of 24 h of water restriction was the one that promoted the highest similarities in the characteristics assessed to those in animals receiving water ad libitum. Therefore, water restriction periods should not exceed 24 h for feedlot animals in situations of severe water shortage.

# 1. Introduction

Water is an essential nutrient for animal production, but its availability is often a limiting factor for livestock in arid and semi-arid regions in the world (Alamer, 2010). In these regions, the animals mostly consume feed with low moisture content, low nutritional value and have irregular and limited access to water.

Sheep and goats can tolerate water shortage by activating some mechanisms to save this nutrient, which reduces the losses and increases the ability to withstand drought (Alamer, 2009). One mechanism to support low water availability is to reduce feed intake to lower the metabolic rate, which works as an adaptation to water conservation, since the animal will generate less heat in the digestive process, reducing the dissipation by evapotranspiration at high ambient temperatures (Maloiy et al., 2008). Furthermore, they tolerate loss of body water > 20% due to the ability of the rumen to store water (Jaber et al. 2004).

According to Barbour et al. (2005), although small ruminants in arid and semi-arid regions can survive up to a week with little or no water, the deficiency of this nutrient adversely affects the homeostasis, body

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weight, reproductive rate and disease resistance, besides the possibility of presenting negative impacts on the meat quality. However, there are few studies evaluating this practice of water management of animals in the Brazilian semi-arid region.

Thus, periods of water restriction alternating with water supply can be an alternative management to minimize the effects of water deficit in feedlot systems, in regions with water shortage. This study evaluates the effect of the water restriction period in Santa Inês crossbred sheep, through the evaluation of carcass traits, non-carcass components, yield of commercial cuts, physical and chemical characteristics and fatty acid profile of the *Longissimus lumborum* muscle.

# 2. Material and methods

# 2.1. Locality, animals and diets

All experimental procedures described in this work were approved by the Ethics Committee of Vale do São Francisco Federal University, with Protocol 0007/161012. The experiment was conducted at the Experimental Station of Caatinga, belonging to the Brazilian



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#### Table 1

Water supply and restriction periods during the experiment for crossbred Santa Inês lambs.

Period		Tratament (hours water restriction)						
	0	24	48	72	-			
Interval without water (days)	0	1	2	3				
Cumulative period with water access (days)	67	34	23	17				
Cumulative period without water access (days)	0	33	44	50				
Total time for data collection (days)	67	67	67	67				

Agricultural Research Corporation - Embrapa, located in the municipality of Petrolina, Pernambuco, Brazil. During the experimental period (May to July 2013), the minimum and maximum average temperatures recorded were 21.97 °C and 31.22 °C, respectively, with relative air humidity of 60.52% and total rainfall of 22.2 mm.

Thirty two Santa Ines crossbred lambs, with initial mean body weight of  $20.7 \pm 2$  kg and eight months of age were randomly assigned to one of four treatments: T1 = without water restriction (daily water supply); T2 = 24 h of restriction and then water supply for 24 h; T3 = 48 h of restriction and then water supply for 24 h and T4 = 72 h of restriction and then water supply for 24 h, as shown in Table 1.

The lambs were housed individually in covered pens  $(1 \times 2 \text{ m})$ , provided with feeders and drinkers. Each animal represented an experimental unit, constituting 8 replicates per treatment. The confinement lasted 77 days, 10 days for adaptation to diet, water supply and facilities and 67 days for data collection.

The diet offered to sheep was the same for all animals, composed of 50% Tifton grass hay and 50% concentrate, consisting of 69.31% ground corn, 29.79% soybean meal and 0.9% mineral, formulated according to the requirements of the National Research Council (NRC, 2007) for daily weight gain of 200 g. Nutrient and fatty acid composition of experimental diet is listed in Table 2. The analyzes to estimate

#### Table 2

Nutrient and	fatty	acid	composition	of	experimental	diet
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Variable	Tifton hay	Concentrate <sup>b</sup>	Diet (50:50) <sup>c</sup>
Nutrient (g/kg DM)			
Dry matter	871.7	863.7	867.7
Organic matter	911.0	970.5	940.7
Ether extract	13.0	24.2	18.7
Crude protein	125.6	204.8	165.2
NDFcp <sup>a</sup>	688.6	97.3	392.9
Acid detergent fiber	398.5	62.3	230.4
Non-fiber carbohydrates	83.4	636.9	360.1
Lignin	90.9	14.5	52.7
Total digestible nutrients	599.3	838.9	719.1
Fatty acids $(g/100 g)^d$			
Myristic (C 14:0)	1.27	0.09	-
Palmitic (C 16:0)	38.76	17.02	-
Margaric (C 17:0)	0.66	0.09	-
Stearic (C 18:0)	2.76	4.65	-
Arachidic (C 20:0)	1.89	0.43	-
Behenic (C 22:0)	2.28	0.22	-
Lignoceric (C 24:0)	2.57	0.22	-
Palmitoleic (C 16:1:c9)	0.57	0.11	-
Heptadecanoic (C 18:1c9)	3.19	28.27	-
Vaccenic (C 18:1c11)	0.91	2.65	-
Petroselinic (C 18:1c12)	0.40	1.53	-
Linoleic C18:2c9c12	12.91	42.81	-
Linolenic C 18:3n3	20.44	1.05	-

<sup>a</sup> Neutral detergent fiber corrected for ash and protein

<sup>b</sup> Was added to concentrate 0.9% of mineral supplement for sheep containing 136 g of sodium/kg

<sup>c</sup> Diet 50:50 (50% hay and 50% concentrate);

<sup>d</sup> %percentage of total fatty acids.

the dry matter, organic matter, ether extract and crude protein were performed according to AOAC (2007). The neutral detergent fiber corrected for ash and protein (FDNcp) and acid detergent fiber were analyzed according to Van Soest, Robertson, and Lewis (1991) and nonfiber carbohydrates and total digestible nutrients according to Sniffen, O'Connor, Van Soest, Fox, and Russell (1992). The extraction of lipids followed the methodology proposed by Hara and Radin (1978) and the transesterification was performed according to Christie (1982). The results were expressed as percentage of total fatty acids.

# 2.2. Dry matter and water intake

The total diet was provided ad libitum, twice a day, at 8:00 h and 16:00 h, with leftovers previously collected and weighed every day to determine daily intake. Samples of the food provided and leftovers were collected weekly, by animal, which were stored in identified plastic bags and stored in a freezer. In these samples, the dry matter (DMI) analyses were carried out to determine the dry matter intake.

The water was supplied in containers of known volume, renewed twice a day and the leftovers measured. The water intake during performance (WIP) was estimated by calculating the difference between the amount of water offered and its surplus, discounting the water lost by evaporation. The WIP (kg) was obtained by multiplying the water consumption by the number of days with access to water during the experimental period (WIP T1 = water consumption × 67 days; WIP T2 = water consumption × 34 days; WIP T3 = water consumption × 23 days, WIP T4 = water consumption × 17 days). To estimate the evaporation, buckets with water were distributed at strategic points in the shed, so that after weighing, this loss (evaportion) was added to the calculation of the WIP per animal.

The water intake via drinker (WID) was estimated by the difference between the amount of water supplied and the surplus and the daily evaporation of water (WID (kg/day) = water supplied – (surplus + evaporation)).

The production of metabolic water (MW) was estimated from the chemical analyses of the diets and calculated by multiplying the consumption of digestible carbohydrate, protein and ether extract by the factors 0.60; 0.42 and 1.10, respectively (Church, 1976; Taylor, Spinage, & Lyman, 1969).

The water use efficiency was estimated by the ratio of water intake to dry matter intake (WI:DMI). Intake of water per kg of carcass produced (WI: kgCAR) was calculated using the relation between the water intake via the drinking fountain and the hot carcass weight.

# 2.3. Slaughter and carcass traits

After the confinement period, the lambs were weighed to obtain the final body weight (BW), and then subjected to solid fasting, receiving only water for 18 h and weighed again to obtain the body weight at slaughter (BWS). Next, the animals were stunned and slaughtered according to protocols established in Regulation of Industrial and Sanitary Inspection of Animal Products - RIISPOA (Brasil, 1997), and subsequently skinned, gutted and removed the head and the extremities of the limbs.

Weights of rumen-reticulum, omasum, abomasum, small and large intestine (full and empty), blood, skin, paws, spleen, liver (without gall bladder), heart, respiratory system and trachea, kidney, head and tongue, reproductive organs (penis and testicles) and empty bladder were also recorded.

Carcasses were weighed to obtain the hot carcass weight (HCW) and to determine the hot carcass yield (HCY). Subsequently, the carcasses were cooled at 4 °C for 24 h, and weighed to obtain the cold carcass weight (CCW) and to calculate the cold carcass yield (CCY), cooling losses of the carcass (CLC) and true carcass yield (TCY) according to Cartaxo et al. (2009). The empty body weight (EBW) was calculated as BWS - gastrointestinal content. After cooling, morphometric measurements were taken on carcasses to calculate the carcass compactness indices (CCI (kg/cm)) = CCW/internal carcass length and leg compactness indices (LCI) = rump width/leg length, according to Souza et al. (2013).

Carcasses were cut lengthwise to obtain the half-carcasses, which were weighed individually. The left half was cut into five anatomical regions: neck, shoulder, rib, loin and leg. In the dorsal portion of *Longissimus lumborum* (LL), at the level of the 13rd thoracic vertebra, measurements of maximum and minimum thickness of subcutaneous fat were taken, using a caliper. To determine the loin eye area (LEA), we used transparent paper to draw the area and then measured the area with the aid of the software Autocad.

# 2.4. Meat quality measurements

Loins of each carcass were identified, conditioned in plastic bags and frozen in a freezer at -18 °C. After 7 days, loins were thawed for 10 h under refrigeration at 10 °C and were dissected to separate the muscle, bone, subcutaneous fat, intermuscular fat and other tissues, which were individually weighed for calculation of yields. Muscle tissue was used to determine the moisture, crude protein and mineral matter contents, according to AOAC (2007) and the ether extract content was determined in an extractor device (ANKOM TX10), according to the methodology proposed by AOCS (2009).

The water holding capacity (WHC) was calculated by the filter paper press method (Hamm, 1986), and the result was expressed in percentage of water exuded compared to the initial sample weight [WHC (%) = [(final weight  $\times$  100)/initial weight]].

To determine the cooking losses (CL), a sample of 50 g in natura Longissimus lumborum muscle was used, free of visible connective tissue. Each sample was divided into 2 subsamples, weighing approximately 25 g and then wrapped in aluminum foil and grilled on plate preheated to 175 °C. The samples were turned over and the internal temperature of the meat was monitored through a digital spit thermometer until it reached 75 °C in the geometric center of the meat. Then, they were cooled at room temperature, taken out of foil and weighed again to calculate the cooking losses.

The following analysis evaluated the shear force (SF), where each sample of grilled meat was cut into two 1cm<sup>2</sup> cubes (totaling 4 replicates for each animal) in the direction of the muscle fibers, and then cut into a texture analyzer (TA-XT Express Texture Analyzer, Stable 165 Micro Systems, Godalming, UK) equipped with a Warner-Bratzler 166 shear force device (2.00 mm/s speed). To classify the meat texture, we adopted the interpretation of Cezar and Sousa (2007).

# 2.5. Fatty acid profile

The composition of the fatty acids present in the lipid extract was obtained using 7 g muscle tissue collected from the *Longissimus lumborum* muscle after dissection (removal of fascia, connective tissue, subcutaneous and intermuscular fat).

The extraction of total lipids from muscle tissue followed the methodology proposed by Hara and Radin (1978), and the transesterification was performed according to Christie (1982). Fatty acid methyl esters (FAME, %) in meat were quantified was carried out by gas chromatography in a Thermo Finnigan Trace-GC Ultra equipment with a flame ionization detector (FID) and a capillary column CP-Sil 88 (Varian), 100 m in length, 0.25  $\mu$ m inner diameter, 0.20  $\mu$ m thick film. Hydrogen was used as the carrier gas at a flow rate of 1.8 ml/min. The initial oven temperature program was 70 °C, 4 min waiting time, 175 °C (13 °C/min) 27 min waiting time, 215 °C (40 °C/min) 9 min waiting time and then increasing 7C°/min to 230 °C, remaining for 5 min, totaling 65 min. The vaporizer temperature was 250 °C and that of the detector 300 °C, according to the temperature program described by Ribeiro et al. (2011).

An aliquot of 1 µl esterified extract was injected into the

chromatograph and the identification of individual fatty acids was performed by comparison of the retention times of the methyl esters presented by the Supelco TM Component FAME Mix chromatography standard (cat 18,919 Supelco, Bellefonte, PA). The fatty acids with cis and trans isomers were separated by chromatographic run along the 100 mm column. No specific treatment was performed for this, and the sample was extracted and esterified as a whole. The concentration of fatty acids was determined by the percentage of the area of a determined fatty acid when added to the areas of all the peaks present in the sample. The results were expressed as g/100 g identified total fatty acid methyl esters.

The nutritional quality of the lipid fraction was evaluated by the composition data of fatty acids, using the calculations of atherogenicity index (AI) = {(C12:0 + (4 × C14:0) + C16:0)}/ $\Sigma$ AGMI +  $\Sigma \omega 6 + \Sigma \omega 3$ ) and thrombogenicity index (TI) = (C14:0 + C16:0 + C18:0)/{(0,5 ×  $\Sigma AGMI$ ) + (0,5 ×  $\Sigma \omega 6$  + (3 ×  $\Sigma \omega 3$ ) + ( $\Sigma \omega 3/\Sigma \omega 6$ )} according to Ulbricht and Southgate (1991). AI indicates the risk of atherosclerosis and TI, the platelet aggregation.

#### 2.6. Statistical analysis

The experiment was a completely randomized design with four treatments and eight replications. The mathematical model used was  $Y_{ij} = \mu + H_j + e_{ij}$ , where  $Y_{ij}$  = value referring to observation of repetition "i" of treatment "j";  $\mu$  = overall mean;  $H_j$  = effect of treatment "j" (0, 24, 48 and 72 h of restriction) and  $e_{ij}$  = random error associated with observation. The half carcass weight was used as covariate in the analysis of the carcass cuts.

The results were analyzed by the software Statistical Analysis System - SAS (version 9.1, 2003), with previous analysis of normality of the residuals by the Shapiro-Wilk test (PROC UNIVARIATE) and the variances compared by orthogonal contrasts (linear, quadratic and deviation from the quadratic model) with a significance level of 5% by PROC GLM. After analysis of contrasts, when significant, we determined the parameters of the regression equations using PROC REG.

# 3. Results and discussion

# 3.1. Water intake and dry matter

The animals that were subjected to water restriction during 72 h showed a reduction of 52.4% (P < 0.05) in the total water intake (WIP) in relation to the animals that received water ad libitum during the experimental period (Table 3). Even observing an increase in daily water intake via drinker (WID) when it was available for animals with 72 h of restriction, the amount of water ingested did not compensate the amount consumed in the ad libitum group. This because the days of consumption were reduced, that is, the animals in the control group ingested water for 67 days, while the animals that underwent water restriction for 72 h only consumed water for 17 days.

The WIP of the animals with 24 h water restriction was 8.1% smaller compared to the control group. The animals with water restriction of 48 h presentend reduction of 34.8% in water intake. This demonstrates the similarity of the total water intake during the performance of the group without restriction with the group of animals with water restriction of 24 h. Ruminant animals, particularly sheep, may survive the dehydration of up to 20%, because of the capacity of the rumen in storing water, which can be used under low availability of this nutrient (Casamassima et al., 2008).

When water was available for the animals with restriction, WID (kg/ day) increased (P < 0.05) according to the increase in water restriction period. The mean consumption for animals that had free access to water was 3.43 kg/day. The sheep with water restriction of 72 h consumed 7.08 kg, that is, 121% daily volume when compared to the animals with free access to drinking water. This is because, in an attempt to quench thirst, the animals ingest more in the first 60 min of access to

#### Table 3

Dry matter and water intake by Santa Ines crossbred lambs submitted to different periods of water restriction.

Variable	Water	on (h)	SEM	$\mathbb{R}^2$	P-value		
	0	24	48	72			Lin.
DMI, kg/day <sup>1</sup> WIP, kg <sup>2</sup> WID, kg/sheep/ day <sup>3</sup>	0.93 229.6 3.43	0.93 211.0 6.21	0.74 149.8 6.51	0.65 120.2 7.08	36.38 10.66 0.32	49.58 59.58 60.00	0.0004 < 0.0001 < 0.0001
WIF, kg/day MW (kg/day) <sup>4</sup> WI:DMI <sup>5</sup> WI/kgCAR <sup>6</sup>	0.12 0.51 3.67 15.69	0.12 0.47 6.68 14.39	0.10 0.38 8.78 12.00	0.09 0.37 10.89 10.92	0.006 0.013 0.13 0.53	- 74.42 31.04 44.69	0.1043 < 0.0001 0.0023 < 0.0001

Dry matter intake (DMI);Water intake during performance (WIP); Water intake via drinker (WID); Water intake via food (WIF); Metabolic water (MW); Water intake: dry matter intake (WI:DMI); Water intake per kg carcass (WI/kg CAR); SEM = standard error of the mean;  $R^2$  = coefficient of determination; Lin: significance for linear effect. Significant at 5% probability; RE = regression equation. RE.<sup>1</sup>:  $\hat{Y} = 0.9647-0.427x$ ; RE.<sup>2</sup>:  $\hat{Y} = 235.456-1.6099x$ ; RE.<sup>3</sup>:  $\hat{Y} = 4.0649 + 0.0473x$ ; RE.<sup>4</sup>:  $\hat{Y} = 0.5125-0.00222x$ ; RE.<sup>5</sup>:  $\hat{Y} = 3.729-0.0137x$ ; RE.<sup>6</sup>:  $\hat{Y} = 15.74143-0.06911x$ .

the water. The animals with water restriction of 24 h consumed 14% more when compared to the animals that received water daily. In addition, when water was given ad libitum again, there was no sign of compensatory increase in food intake.

The extending the period of water restriction reduced the dry matter intake by 30% (P < 0.05), of the group without water restriction compared to the one that with water restriction of 72h (0.93 and 0.65 kg/animal/day, respectively), inferring that, because of the unavailability of water, animals reduce this consumption. Due to periods of water unavailability, in the attempt to quench thirst, the animals ingested more water per day at the time of the supply, mainly the animals that spent three days without ingesting water, even consuming more than the animals with free access to water. This may have caused ruminal fill and, consequently, hypertonicity of the rumen-reticulum musculature caused by the accumulation of ingested water, reducing DMI. An abnormal prandial increase in ruminal fluid osmolality contributes to the suppression or reduction of food intake during water restriction (Burgos, Langhans, & Senn, 2000). Despite a marked decrease in DMI of animals subjected to 48 and 72 hour-restriction, the DMI of animals with 24 h of water restriction was close to the intake of animals with daily access to water.

According to Alamer & Al-hozab, 2004), the reduction in dry matter intake due to water restriction, is an adaptation mechanism to reduce costs related to the use of water in food digestion process, thus resulting in greater water conservation. This concomitantly reduces heat production (metabolism) and increases the water retention, sufficient to achieve a new equilibrium over a longer period of water restriction. Abioja, Osinowo, Adebambo, Bello, and Abiona (2010) explained that the reduction in dry matter intake from the water restriction can be the result of the need of water for moistening the bolus and transportation of the gastrointestinal tract content.

Despite the reduction in dry matter intake, there was no effect (P > 0.05) of the water restriction on water intake by food (WIF). However, there was a decreasing linear effect (P < 0.05) for MW (kg/day). Misra and Singh (2002) found no effect on metabolic water intake when subjected goats in semi-arid regions of India to 48 h of water restriction. Metabolic water is produced during the oxidation of the hydrogens contained in the main nutrients, with 1 g of protein, carbohydrate and fat producing 0.42 g; 0.60 g and 1.10 g water for each nutrient, respectively (Church, 1976). With the reduction in DMI, the availability of nutrients for oxidation also reduced, then, there was a decrease of the water produced during the catabolism.

According to the NRC (2007), for each kilogram of dry matter

#### Table 4

Weight,	, yield a	and indi	ices of	f Santa	Ines	crossbred	lambs	submitted	to	different
periods	of wat	er restri	ction.							

Variable	Water re	striction (	h)		SEM	$\mathbb{R}^2$	P-value
	0	24	48	72			Lin
BWS, kg <sup>1</sup>	32.62	32.29	26.77	25.88	0.89	66.80	0.0011
EBW, kg <sup>2</sup>	25.33	25.04	21.90	20.08	0.69	49.43	0.0015
HCW, kg <sup>3</sup>	14.90	14.26	11.27	10.85	0.47	43.48	0.0051
CCW, kg <sup>4</sup>	14.32	13.76	10.76	10.45	0.46	42.87	0.0048
HCY, % <sup>5</sup>	45.69	44.17	42.10	41.91	0.48	53.26	0.0007
CCY, % <sup>6</sup>	43.91	42.62	40.23	40.38	0.46	54.65	0.0005
CCR, %	4.10	3.83	4.11	4.84	0.11	-	0.9848
TCY, %	54.89	56.19	56.62	50.10	0.45	-	0.6430
CCI, kg/cm <sup>7</sup>	0.23	0.24	0.21	0.20	0.005	73.28	0.0116
LCI	0.62	0.60	0.62	0.62	0.008	-	0.7941

BWS: body weight at slaughter; EBW: empty body weight; HCW: hot carcass weight; CCW: cold carcass weight; HCY: hot carcass yield; CCY: cold carcass yield; CCR: carcass cooling rate; TCY: true carcass yield; CCI: carcass compactness index; LCI: leg compactness index; SEM = standard error of the mean;  $R^2$  = coefficient of determination; Lin: significance for linear effect. Significant at 5% probability; RE = regression equation. RE.<sup>1</sup>:  $\hat{Y} = 33.78782-0.1140 \times$ ; RE.<sup>2</sup>:  $\hat{Y} = 26.93913-0.09924x$ ; RE.<sup>3</sup>:  $\hat{Y} = 14.99189-0.0563x$ ; RE.<sup>4</sup>:  $\hat{Y} = 14.42074.0-0.05427x$ ; RE.<sup>5</sup>:  $\hat{Y} = 45.6435-0.0563x$ ; RE.<sup>6</sup>:  $\hat{Y} = 44.007-0.0552x$ ; RE.<sup>7</sup>: 0.24923-0.00089744x.

ingested, the animal should consume 2.87 l of water. All animals had a higher intake than that recommended by the NRC (2007). According to this study, there was an increase (P < 0.05) in the ratio WI: DMI when the animals were subjected to different periods of water restriction, because, as the restriction periods increased, the animals consumed more water when it was available. The WI: DMI ratio of the animals that underwent water restriction of 24 h, 48 h and 72 h were 182.0, 239.2 and 296.7%, respectively, higher than the intake of animals that received water daily.

# 3.2. Carcass traits

The body weight at slaughter (BWS) decreased linearly (P < 0.05) with increasing periods of water restriction (Table 4). Studies on water restriction indicate that a part of the body weight reduction in ruminants is related to the combined effect of body water loss and decreased feed intake (Alamer, 2006, 2009; Hamadeh et al., 2006; Silanikove, 1992) or mobilization of fat for energy production to compensate for the lower feed intake (Jaber et al., 2004) during the water restriction periods. During this period, the body weight of the animals is reduced, also decreasing the requirement of nutrients. The BWS of the animals subjected to water restriction, demonstrating that the deprivation of water for 24 h followed by 24 h of hydration did not affect the final weight of the animals.

The EBW, HCW and CCW decreased (P < 0.05) with increasing water restriction period. Prolonged periods of water restriction reduce the gastrointestinal passage rate, reducing body weight at slaughter and, consequently, empty body weight and hot carcass weight. The reduction in nutrient intake for animals may explain the observed decrease in body weight at slaughter, empty body weight, hot carcass weight and cold carcass weight, since these parameters reflect directly on the animal performance and are natural indicators of the ingestion of nutrient (Urbano et al., 2013). In the dry season, when food and water availability is limited, regression of productive indexes and animal performance may occur, reducing productivity (Ponnampalam et al., 2018). Tibin, Bushara, Elemam, Tibin, and Jadalla (2012) evaluated the sheep carcass traits in the desert, which were subjected to water supplies with intervals every 2-3 days, water ad libitum with and without supplementation in the diet, and found lower body weight at slaughter, hot carcass weight, half carcass and empty body weight for animals that

had restricted access to water, corroborating data from this study.

In addition, the HCY and CCY decreased (P < 0.05) with increasing water restriction periods. This reduction can be explained by the decrease in BWS, HCW and CCW, due to reduction in nutrient supply. Although the observed decrease in HCY e CCY, the periods of the water restriction did not influence (P > 0.05) the TCY. Although the reduction of BWS, HCW, CCW, HCY e CCY with the water restriction, the group of animal with restriction every 24 h showed similar weights the group that was not submitted the water restriction.

The periods of the water restriction did not influence (P > 0.05) the cooling losses (CCR), with a mean value of 4%. These losses can vary from 3.0 to 4.0%, according to the uniformity of fat, sex, weight and chilling temperature (Reis et al., 2001). The CCR can be indicative of adequate fat cover and protection against cooling in the refrigeration chamber, thus avoiding shortening by cold and excessive loss of water from the meat (Safari, Fogarty, Ferrier, Hopkins, & Gilmour, 2001).

The water restriction did not influence (P > 0.05) the TCY, with a mean value of 54%. According to Sañudo and Sierra (1986), carcass yields range from 40 to 60%, according to breed, crosses and breeding system. Therefore, the data obtained in this study are consistent with those described by these authors.

There was a reduction (P < 0.05) of the CCI as the water restriction periods increased. Compactness indices indicate the amount and/or storage capacity of meat in the carcass and leg, which decreased with increasing restriction, i.e. the carcasses had reduced capacity of storing tissues, which agrees with the weights of retail cuts. However, LCI was not affected (P > 0.05) by the water restriction periods.

There was a decrease (P < 0.05) in the ratio of water intake per kilogram carcass (IW/kg CAR), that is, the animals of the control group ingested 229.621 of water throughout the experimental period, to produce 13.88 kg carcass, while animals with 72 h of water restriction ingested 29.63 L to produce 10.04 kg, i.e., less 199.991 of water, with difference of 3.84 kg carcass. The carcass production of the animals with water restriction for 72 h was similar to that of the animals receiving water every day, saving 2001 of water, which would be feasible under conditions of water shortage for feedlot animals. With the increase of the water restriction periods, there was reduction in the dry matter intake and decrease in the weight at slaughter, however, it did not affect the deposition of fat in the carcass. Thus, it is suggested that water restriction reduces the energy metabolism to conserve water and compensate for the reduction in food intake. When water supply is limited, there is a close interrelation between the amount of dry matter consumed and the amount of water ingested, which is a consequence of the low relationship between energy and water intake (Silanikove, 1989).

#### 3.3. Carcass cuts and tissue composition

The water restriction periods did not affect the HCW and the carcass cut weights (P > 0.05). However, when HCW was used as a covariate, there was a significant effect on carcass cut weights (Table 5).

The proportion of muscle did not differ (P > 0.05) as shown in Table 6. This is related to the order of growth (estimated by allometry coefficients) of the muscle, which has an isometric development and evolves parallel to the growth of the carcass. These results confirm the anatomical harmony law of Boccard and Dumont (1960), according to which the relative proportions of the different body regions are similar in carcasses of similar weight and fattening.

The water restriction did not affect (P > 0.05) the proportion of subcutaneous fat, intermuscular fat, minimum (FTMin) and maximum (FTMax) fat thickness in relation to the loin. It was expected that the animals in the control group had the highest percentage of fat, since they have higher BWS and fat had a late development, but the water restriction did not affect this parameter. It has been reported that under a low nutritional level, the priority of nutrient supply for each part of the body depends on its rate of development and metabolic rate (Atti,

#### Table 5

Weight of carcass cuts of Santa Ines crossbred lambs submitted to different periods of water restriction.

Variable	Water	restrictio	on (h)		SEM	$\mathbb{R}^2$	P-value	
(kg)	0	24	48	72			HOURS	$\rm HCM^1$
HCW Leg <sup>2</sup> Loin <sup>3</sup> Rib <sup>4</sup> Shoulder <sup>5</sup> Neck <sup>6</sup>	7.11 2.31 0.62 1.92 1.28 0.56	7.05 2.29 0.65 1.94 1.27 0.60	6.18 2.03 0.49 1.53 1.13 0.53	5.65 1.94 0.49 1.48 1.08 0.46	0.22 0.07 0.03 0.07 0.03 0.02	72.57 66.23 64.58 72.64 80.70 54.38	NS NS NS NS NS	0.0001 0.0001 0.0001 0.0001 0.0001 0.0001

HCW = half carcass weight; SEM = standard error of the mean; Significant at 5% probability; 1HCM: half carcass weight as a covariate; NS: not significant.

# Table 6

Tissue composition, loin measurements and physical and chemical characteristics of meat from Santa Ines crossbred lambs submitted to different periods of water restriction.

Variable	Water	restrictio	n (h)	SEM	$\mathbb{R}^2$	P-value	
	0	24	48	72			Lin
Muscle, %	49.81	47.58	53.11	54.35	1.05	-	0.1032
Subcutaneous fat, %	10.88	9.88	11.17	8.92	0.56	-	0.3606
Intermuscular fat, %	3.77	3.50	3.95	3.71	0.26	-	0.9159
Bone, %	22.26	26.79	17.55	21.36	0.99	-	0.1170
Others, %	11.21	11.73	11.38	10.75	0.31	-	0.5423
FTMin, mm	1.58	1.18	1.18	1.16	0.13	-	0.2774
FTMax, mm	5.54	5.36	5.27	5.21	0.33	-	0.7290
Loin eye area, cm <sup>2</sup>	10.01	8.12	8.43	8.98	0.29	-	0.4897
WHC, %	68.91	66.79	66.23	65.99	0.64	-	0.1066
CL, %	27.96	30.65	30.03	30.72	0.65	-	0.1429
SF (N/cm <sup>2</sup> ) <sup>1</sup>	30.50	30.79	38.15	45.79	0.25	60.29	0.0055
Water, %	75.00	75.00	76.00	75.00	0.39	-	0.8492
Crude protein, %	20.89	21.01	20.08	21.41	0.23	-	0.7589
Ether extract, %	2.31	3.80	2.00	1.97	0.41	-	0.4451
Mineral matter, %	1.12	1.19	1.13	1.11	0.02	-	0.6181

FTMin = minimum fat thickness; FTMax = maximum fat thickness; WHC = water holding capacity; CL = cooking losses; SF = Warner-Bratzler shear force; SEM = standard error of the mean;  $R^2$  = coefficient of determination; RE.<sup>1</sup>:  $\hat{Y} = 28.52 + 0.0222x$ , Lin: significance for linear effect. Significant at 5% probability.

# Mahouachi, & Rouissi, 2006; Kamalzadeh, Koops, Van Bruchem, & Brangma, 1998).

However, there was no effect (P > 0.05) for bone percentage in the loin, with a mean value of 22%. Bone is an early maturing tissue and at 8 months (growing animals) is less affected by diet. For tissue composition, muscle yield was superior to other tissues. Possibly, the highest percentage of muscle over fat and bones, in the analyzed cut, was influenced by the age of the animals, which were young (eight months).

There was no effect (P > 0.05) for loin eye area with water restriction, with a mean value of  $8.93 \,\mathrm{cm}^2$ . The *Longissimus lumborum* muscle has late development, thus, the young age of the animals would justify the values found. This result was not to expected, once the reduction in water consumption decreases DM intake and, consequently, the supply of nutrients, mainly of protein for the muscular development.

# 3.4. Meat physical and chemical parameters

The water holding capacity (WHC) was not influenced by the water restriction, with a mean value of 67% (Table 6). The WHC of meat is the ability of retaining water during the application of external forces and affects the juiciness at the time of consumption (Henchion et al. 2014). In addition, it has great importance in storage and shelf life of the product. Possibly, the water restriction of up to 72 h did not affect the

pH of the postmortem muscle, and consequently, the isoelectric point of proteins (Munasinghe & Sakai, 2004) maintaining the meat ability to hold water.

The water restriction periods did not influence (P > 0.05) the cooking losses, with a mean value of 29.84%. However, SF increased according to the extended period of water restriction. Meat of the animals with water restriction for 24 h had SF of 30.8 N/cm<sup>2</sup>, classified as medium tenderness (22.4 to 35.6 N/cm<sup>2</sup>). The water restriction for 48 and 72 h resulted in meat with SF of 38.1 N/cm<sup>2</sup> and 45.8 N/cm<sup>2</sup>, respectively, classified as hard (35.7 to 53.4 N/cm<sup>2</sup>), following the scale suggested by Cezar and Sousa (2007).

These results can be explained by the fact that animals that were subjected to a longer period of water restriction suffered from preslaughter stress. In addition, there was a reduction in food intake, which have reduced glycogen storage in the muscle. In stress conditions, there is an increased glycolytic activity, rapid protein denaturation, accelerating the rigor mortis process, making the meat of the animals tougher (Carragher & Matthews, 1996). No other published study to our knowledge provides a comparison on the effect of water restriction on meat tenderness.

The determination of water content in meat is one of the most important measures used in food analysis, since it is related to its composition, stability and quality, and can influence the storage, packaging and processing (Jiménez Colmenero, 1996), moreover exerts influence on carcass dressing. Nevertheless, the water content in the LL muscle was not influence (P > 0.05) by water restriction, with mean value of 75%, corroborating Madruga et al. (2008), which stated that the sheep meat has about 73% moisture.

The contents of crude protein, fat and minerals in meat were not affected (P > 0.05) by water restriction with mean values of 21; 2 and 1%, respectively. According to Zeola, Silva Sobrinho, Gonzaga Neto, and Marques (2004), the average protein in sheep meat is 19%, ether extract is 2% and mineral matter is 1%, values similar to those found herein. Thus, different periods of water restriction had no influence on the chemical composition of LL muscle of animals.

# 3.5. Non-carcass components

The water restriction did not influence the head weight (P < 0.05), suggesting that the growth pattern is related to the coefficient of allometry, in which the bone growth is intermediate (Rosa, Pires, Silva, & Motta, 2002), occurring in the initial growth phase and is less affected by diet (Table 7).

The weights of the paws linearly decreased (P < 0.05) with increasing of water restriction period. Despite having an early development, the weight of paws reduced possibly due to the decrease the dry matter intake, due to the reduced interval of water supply, inferring the need for nutritional support for development.

The rumen-reticulum weight decreased (P < 0.05) with longer periods of water restriciton, which is explained by the reduction of DMI and water, which may have caused less developed and distention of the organ. Kremer, Lorenzi, and Barbato (1989) also mentioned that the development of the rumen-reticulum is related to the animal weight, corroborating this study. Therefore, animals with longer interval of water restriction had lower body weight at slaughter, which reinforces the fact that the lack of water has caused lower food intake, the development and the distension of these organs.

Water supply intervals did not affect (P > 0.05) the weight of omasum, abomasum, small and large intestine, blood, spleen, heart, head and tongue and bladder. Drouillard et al. (1991) reported that these organs of importance to the animal are proportionally larger at birth and, consequently, develop less in postnatal life.

The skin, due to the good softness and elasticity, is the most important non-carcass component, reaching 10% of the animal value; however, there was a reduction (P < 0.05) in weight by 23%. The reproductive organs (RO) decreased linearly (P < 0.05) by 24 and

#### Table 7

Non-carcass components of Santa Ines crossbred lambs submitted to different periods of water restriction.

Variable	Water restriction (h)			SEM	$\mathbb{R}^2$	P-value		
(kg)	0	24	48	72			Lin	Quad
RR <sup>1</sup>	0.76	0.78	0.69	0.61	0.02	42.06	0.0016	0.1432
Omasum	0.09	0.13	0.07	0.19	0.03	-	0.3936	0.5250
Abomasum	0.15	0.17	0.15	0.16	0.007	-	0.9799	0.6359
SI	0.82	0.82	0.76	0.77	0.03	-	0.5112	0.9404
LI	0.23	0.22	0.18	0.21	0.01	-	0.3376	0.2872
Blood	0.95	1.07	0.78	0.79	0.04	-	0.0621	0.5450
Skin <sup>2</sup>	2.70	2.80	2.54	2.08	0.09	42.41	0.0058	0.0946
Spleen	0.05	0.06	0.05	0.04	0.002	-	0.1556	0.0957
Liver <sup>3</sup>	0.53	0.49	0.44	0.40	0.02	65.30	0.0033	0.9783
Heart	0.12	0.12	0.11	0.11	0.04	-	0.1628	0.6959
RSTE <sup>4</sup>	0.57	0.57	0.51	0.44	0.01	47.95	0.0032	0.3078
Kidneys <sup>5</sup>	0.08	0.09	0.08	0.07	0.02	52.22	0.0770	0.0002
H + T	1.51	1.55	1.43	1.38	0.04	-	0.1717	0.5684
Paws <sup>6</sup>	0.74	0.75	0.66	0.57	0.02	55.31	< 0.001	0.3906
RO <sup>7</sup>	0.37	0.33	0.27	0.28	0.01	24.69	0.0161	0.4465
Bladder	0.02	0.01	0.01	0.01	0.002	-	0.2293	0.2918

RR = rumen-reticulum; SI = smallintestine; LI = largeintestine: RSTE = respiratory system, trachea and esophagus; H + T = head and tongue; RO = reproductive organs (penis and testicles); SEM = standard error of the mean;  $R^2$  = coefficient of determination; Lin: significance for linear effect. Quad: significance for quadratic effect. Significant at 5% probability; RE.<sup>1</sup>:  $\hat{\mathbf{Y}} = 0.7957 - 0.0023 \mathbf{x};$ RE = regressionequation: ER.2: ER.<sup>3</sup>:  $\hat{Y} = 0.55234 - 0.00237x$ ; ER.<sup>4</sup>:  $\hat{Y} = 0.58879$  $\hat{\mathbf{Y}} = 2.7766 - 0.01125 \mathbf{x};$ ER.<sup>5</sup>:  $Y = 0.07735 + 0.000608x - 0.0000099x^{2}$ ; -0.0025x: ER 6:  $\hat{Y} = 0.7737 - 0.00278x$ ; ER.<sup>7</sup>:  $\hat{Y} = 0.3661 - 0.1453x$ .

28%, respectively. Both the skin and the reproductive organs presented isogonic growth, that is, the growth rate was similar to EBW.

There was a 23% (P < 0.05) reduction in the weight of respiratory system and trachea (RSTE). During periods of water shortage, some physiological mechanisms are activated, such as reduction of respiratory rate, resulting in decreased water losses (Alamer, 2009), as a way to save this nutrient. The weight of the RSTE was possibly reduced as the water restriction period increased in response to the low water intake and consequently the lower respiratory rate in an attempt to reduce the water removed during respiration.

The increase in the period of water restriction promoted a reduction (P < 0.05) in liver weight, which is important for various metabolic processes, especially for energy and protein metabolism, reducing 17% at 72 h of water restriction compared to daily supply. Liver and kidneys are priority organs in animal metabolism and the reduction in liver may be indicative of the reduction of metabolic rate, since the water consumption promoted reduction in dry matter intake (Camilo et al., 2012), with a notable atrophy for receiving feed below the maintenance level.

There was a quadratic effect (P < 0.05) for the weight of the kidneys of sheep, with the highest value for lambs subjected to 24 h of water restriction. When there is water deficit, there is stimulation of antidiuretic hormone (ADH) secretion, which increases water reabsorption in renal collecting tubules. This mechanism also activates the thirst center, increasing water intake, increasing activity on renal flow and thus stimulating the growth, providing greater weight (Naves, Vilar, Costa, Domingues, & Casulari, 2003).

The water restriction period did not influence (P > 0.05) the volume of the blood of animals, even with restriction of 72 h, this must have occurred because the initial response of the body to the negative balance of water is the retention of body fluids (Kaliber, Koluman, & Silanikove, 2015), maintaining constant blood volume of the animals. It is emphasized the importance of studies on these components, because some of them serve as food for the human population, such as the head, liver, heart, kidneys, lungs and digestive tract (Osório, Oliveira, Osório,

#### Table 8

Fatty acids of *Longissimus lumborum* muscle (% of total fatty acids) of lambs submitted to different periods of water restriction.

Variable	Water	restrictio	n (h)	SEM	$\mathbb{R}^2$	P-value	
	0	24	48	72			Lin
ΣSFA	43.87	43.93	44.80	45.98	0.47	_	0.0810
Caproic (C10:0)	0.35	0.36	0.26	0.36	0.019	-	0.6211
Lauric (C12:0)	0.16	0.16	0.12	0.14	0.009	-	0.1958
Myristic (C14:0)	2.00	2.00	2.00	2.00	0.054	-	0.8938
Pentadecanoic (C15:0)	0.25	0.26	0.27	0.25	0.006	-	0.7891
Palmitic (C16:0)	23.00	22.00	23.00	23.00	0.356	-	0.8562
Margaric (C17:0) <sup>1</sup>	0.59	0.65	0.70	0.70	0.012	45.35	0.0008
Isomargaric (C17:0iso)	0.23	0.22	0.21	0.23	0.014	-	0.9766
Stearic (C18:0)	17.00	18.00	18.00	19.00	0.418	_	0.0731
Behenic (C22:0)	0.15	0.14	0.11	0.14	0.012	-	0.6556
ΣUFA	46.36	46.39	46.38	44.35	0.47	-	0.0810
Palmitoleic (C16:1c9)	2.00	2.00	2.00	2.00	0.097	-	0.1341
Oleic (C18:1c9)	41.00	41.00	41.00	38.00	0.782	_	0.1287
Vaccenic	2.00	2.00	2.00	3.00	0.169	51.38	0.0006
$(C18:1 t11)^2$							
Petroselinic	1.00	1.00	1.00	1.00	0.073		0.0754
(C18:1c12)							
ΣPUFA	8.86	7.78	7.76	7.79	0.62	-	0.8583
Linoleic	5.00	5.00	5.00	5.00	0.353	-	0.9038
(C18:2c9c12)							
CLA (C18:2c9t11) <sup>3</sup>	0.27	0.21	0.19	0.15	0.018	63.11	0.0002
Linolenic	0.33	0.28	0.27	0.29	0.018	-	0.5391
(C18:3n3)							
AA (C20:4 n6)	3.00	2.00	2.00	2.00	0.274	-	0.7125
EPA (C20:5 n3) <sup>4</sup>	0.23	0.26	0.28	0.34	0.011	42.18	< 0.0001
DHA (C22:5) <sup>5</sup>	0.03	0.03	0.02	0.01	0.001	52.80	< 0.0001
SFA:UFA	0.75	0.79	0.82	0.82	0.02	-	0.0935
ω6	5.02	4.68	4.35	5.07	0.31	-	0.8144
ω3	0.32	0.28	0.26	0.29	0.17	-	0.5402
ω6:n3	15.60	16.93	16.68	17.43	0.52	-	0.1038
AI	0.54	0.56	0.56	0.56	0.01	-	0.7134
TI	1.40	1.47	1.53	1.53	0.03	-	0.0832
h:H	2.03	2.01	1.96	1.92	0.04	-	0.3496

CLA (conjugated linolenic acid); AA (Aracdonic acid); EPA (eicosapentaenoic acid); C22:5 (docosapentaenoic acid);  $\Sigma$ SFA: sum of saturated fatty acids;  $\Sigma$ MUFA: sum of monounsaturated fatty acids;  $\Sigma$ UFA: sum of unsaturated fatty acids;  $\Sigma$ UFA: sum of unsaturated fatty acids;  $\Sigma$ UFA: sum of unsaturated fatty acids;  $\omega$ : omega 6;  $\omega$ 3: omega 3; AI: atherogenicity index; TI: thrombogenicity index; DFA: desirable fatty acids; h:H: hypocholesterolemic: hypercholesterolemic fatty acids ratio SEM = standard error of the mean; R<sup>2</sup> = coefficient of determination; Lin: significance for linear effect. Significant at 5% probability; Regression equation (RE)<sup>1</sup>:  $\hat{Y} = 0.619 + 0.00095x$ . RE<sup>2</sup>:  $\hat{Y} = 1.684 + 0.0222x$ ; RE.<sup>3</sup>:  $\hat{Y} = 0.0326 - 0.002283x$ .

# Jardim, & Pimentel, 2002).

# 3.6. Fatty acid profile

Regarding the profile of saturated fatty acids of meat, the water restriction period did not affect (P > 0.05) the percentage capric (C10:0), lauric (C12:0), myristic (C14:0), pentadecanoic (C15:0), palmitic (C16:0), stearic (C18:0) and behenic (C22:0) acids (Table 8).

The palmitic (C16:0) and stearic (C18:0) acids were the most prevalent in the lipid profile of meat as to total saturated fatty acids. This was due to the higher concentration of these fatty acids in the diet. The medium chain fatty acids such as myristic (C14:0) and palmitic (C16:0), considered hypercholesterolemic can increase both the total cholesterol concentration in the plasma and raise low density lipoprotein (LDL). While larger chain fatty acids such as stearic (C18:0), they are considered to be neutral for plasma cholesterol (Scollan et al., 2001).

No effect (P > 0.05) of water restriction was detected on the percentage of monounsaturated fatty acids, with mean values of 39 and 1% of C18:1c9 and C18: 1 c12, respectively. Among the unsaturated, it is verified a greater amount of C18:1c9 acid, which varies between 30 and 43% (Sañudo et al., 2000), confirming this study. Oleic acid, recognized for the hypocholesterolemic effect, was the monounsaturated fatty acid with the highest percentage, as observed by Sañudo et al. (2000), who mentioned that this acid ranges from 30 to 43% in lipid profile of the meat.

The increase in the water restriction period provided an increase (P < 0.05) in the vaccenic acid (C18:1 trans11). The water restriction decreased dry matter intake and possibly reduced the population of bacteria responsible for ruminal biohydrogenation. Thus, there was a reduction in the conversion of unsaturated fatty acids, from the diet, into saturated, increasing the concentration of vaccenic acid, intermediate of biohydrogenation, in the rumen. After absorption by the small intestine of the animal, C18: 1 trans11 is deposited in muscle tissue (Van Nevel & Demeyer, 1996). Vaccenic acid is an important precursor in the intermediary metabolism of conjugated linoleic acid (CLA), responsible for approximately 90% CLA of the intramuscular fat of the meat (Nuernberg et al. 2002) Thus, it is expected that the increase in the vaccenic acid content will increase the amount of CLA in the meat, which was not verified in the present study.

The increase in the water restriction period resulted in a reduction (P < 0.05) of CLA in sheep meat. Vaccenic acid is transformed into CLA in ruminant muscle tissue through the action of the enzyme delta-9-desaturase (Smith, Gill, Lunt, & Brooks, 2009). Thus, reductions in water and dry matter, possibly have inhibited the messenger RNA synthesis of the delta-9 desaturase enzyme (Baumgard, Sangster, & Bauman, 2001; Park et al., 2000) and, therefore, the conjugated linoleic acid, decreasing its abundance in the muscle tissue of animals with less access to water. This is not satisfactory in view of the beneficial properties of this acid due to nutraceutical properties. In addition, this fatty acid presents anticancer and beneficial effects to cardiovascular health (Tapiero, Nguyen Ba, & Couvreur, 2002).

EPA and DHA have several cellular functions, are precursors of eicosanoids (prostaglandins, thromboxanes and leukotrienes) that have great importance in cardiovascular health. Moreover, they are essential for maintaining the integrity of the cell membrane and are important mediators of gene expression (Clarke, 2001). There was an increasing linear effect (P < 0.05) of EPA with the water restriction and a decreasing linear effect (P < 0.05) of DHA with the increase in water restriction. The lowest values were found in the meat of animals subjected to 72h of restriction. EPA is formed by desaturation and stretching of  $\alpha$ -linolenic acid (Smith, 2007), so the reduction in feed intake caused by the reduction in water intake probably induced the synthesis of the enzyme delta 6 desaturase and elongases required for conversion of alpha-linolenic into its long chain derivatives (Pawlosky et al., 2003). Increasing EPA incorporation into cell membrane phospholipids results in increased production of eicosanoids that have antiinflammatory characteristics (Calder, 2006). Despite the increasing linear effect for EPA, DHA was reduced in the meat of animals subjected to 48 and 72 h of restriction, indicating that after this restriction period, there was probably a reduction in the synthesis of enzymes required for the conversion of linolenic acid into DHA, as well as specific transport mechanisms for these fatty acids in muscle tissue. DHA is the most important for proper cell membrane function and is vital for the development of the brain and retina (Ramakrishnan et al., 2010).

There was no effect for the sum of saturated (44.14%), unsaturated (55.85%), monounsaturated (47.46%), polyunsaturated (8.39%) fatty acids and SFA: UFA ratio. Dietary fatty acids are affected by rumen microorganisms, especially with respect to polyunsaturated fatty acids, with effects on the content and composition of fatty acids in the muscle. However, the water restriction up to 72 h did not affect the sum of these fatty acids. The  $\omega 6/\omega 3$  ratio was not affected (P > 0.05) by the water restriction period up to 72 h. The SFA: UFA and  $\omega 6: \omega 3$  ratios are used to evaluate the nutritional value of oils and fats and to indicate the cholesterolemic potential. There was no effect (P > 0.05) for AI, IT and

the h: H ratio. These results corroborate the effect reported in this study for the muscle ether extract (Table 6), which showed no interaction with the water restriction.

#### 4. Conclusions

Water restriction up to 72 h reduces water intake and dry matter intake, and consequently body weight and carcass yield. However, the final body weight and carcass yield of the animals subjected to 24 h of restriction was similar to the weight of the animals that received water daily. The weight of the meat cuts was not affected by water deprivation, as well as the tissue and chemical composition of the Longissimus lumborum muscle. The weight of non-carcass components commercially valued as the rumen-reticulum, skin and liver reduced with the water restriction, however, the weights of these were similar between the 24h group and the group with free access to water deprivation. The highest weight of the kidneys of lambs was observed in animals subjected to 24 h of water restriction. The amount of fatty acids in the meat presented little variation due to the water restriction periods, however, there was an increase in EPA and a reduction in CLA and DHA, important for human health. The indices of atherogenicity and thrombogenicity were not affected by water restriction.

This study demonstrates that is possible produce carcass and meat with quality through feedlot lamb in semi-arid region, during the dry season, where do you have severe water shortage, using water restriction intervals up to 24 h. The interval of 24 h of water restriction promoted the highest similarities in the characteristics assessed compared to those in animals receiving water ad libitum. Thus, water restriction periods should not exceed 24 h and may be used to reduce water intake by feedlot sheep in situations extreme water shortage.

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