EFFECT OF UREATED SUGAR BEET PULP ON LACTATING GOAT PERFORMANCE Hend A. Aziz¹ and A.M. Kholif² ¹Animal Nutrition Department, Desert Research Center, Cairo, Egypt. ²Dairy Science Department, National Research Centre, Dokki, Giza, Egypt.

ABSTRACT

Forty female Barki goat does were used to investigate the effect of inclusion of sugar beet pulp (SBP) treated with urea at levels of 2, 4 and 6% urea to replace about 30% of common concentrate feed mixture in the diet of goats on lactating goat performance, milk yield and composition, birth and weaning weight. Five experimental diets were formulated: T1 (control): concentrate feed mixture (CFM)+ berseem hay (BH), T2: CFM + untreated SBP+BH, T3: CFM+SBP treated with 2% urea on DM basis+ BH, T4: CFM + SBP treated with 4% urea on DM basis+ BH and T5: CFM + SBP treated with 6% urea on DM basis+ BH. The experiment feeding period lasted for 140 days. At the end of lactation stage a digestibility trial was carried out to study the effect of the experimental diets on digestibility coefficients, rumen parameters, microbial protein, protozoal count, and blood constituents. The obtained results revealed that T5, T4 and T3 increased (P≤0.05) DM, OM, EE, NFE and CP contents, and decreased ash, CF, NDF, ADF, DAL, cellulose and hemicellulose contents as compared to T2 and T1. Urea treatments of SBP increased (P≤0.05) body weight and feed intake. T4 and T5 had the highest (P≤0.05) DM, OM, EE, CP, CF, NFE, NDF, ADF, ADL, cellulose and hemicellulose digestibility coefficients as well as the highest (P≤0.05) TDN and DCP intakes. T4, T5 and T3 improved (P<0.05) nitrogen balance. water balance, rumen fermentations, rumen protozoal count, milk yield and composition, birth and weaning weight. The present study suggests the replacement of dried sugar beet pulp treated with 4% urea by a part of common concentrate feed mixture of lactating goats.

Keywords: Sugar beet pulp, urea treatment, lactating goats, blood, kid performance.

INTRODUCTION

In Egypt there is a gap in animal requirements and the available feeds, so there is an urgent need to search for more available nonconventional and cheaper feed sources, particularly agricultural by-products as sugar beet pulp (SBP). The major limitations of using these agricultural residues as feed are its low palatability, low digestibility, low protein and high fiber content (El-Ashry *et al.* 2003). SBP is a solid vegetable matter that remains after sugar extraction from sliced sugar beets, which has been estimated to comprise 6% dry matter (DM) of the weight of beet root (Talha *et al.* 2002). This by-product has a high content of crude fiber (CF, 17–22%) and low content of crude protein (CP, 8-11%) as reported by Papadomichelakis *et al.* (2004). Although its crude fiber content (El-Ashry *et al.* 2000). Consequently, SBP was considered as an energy source in mixed rations of dairy cattle and sheep (El-Badawi *et al.* 2003; Sherien 2005).

Large areas of several developing countries are cultivated by sugar beets. Therefore, there are huge amounts of SBP without beneficial usage

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and might be considered as a byproduct. However, if these poor quality byproducts were utilized by improving their digestibility and nutritive value, it might solve part of feed-shortage problems in such countries. Chemical and biological treatments have been tested by various researchers to increase the nutritive value, palatability, DM intake, protein content, and the digestibility of poor quality roughages (Aziz *et al.* 2008; Omer *et al.* 2012). Several chemical treatments including urea treatments aimed to disrupt the lignin carbohydrate complex which has been tested in attempts to improve the accessibility of structural carbohydrates to celluloytic microorganisms; chemical pretreatment appeared to improve more particularly the digestibility of the hemicellulose fraction (Aziz 2009).

The aim of this study was to improve the nutrient value of SBP using chemical treatments (2, 4 and 6% urea) to replace a part of concentrates. Also, the effect of treatments on digestibility, rumen fermentations, ruminal protozoa count, milk yield and composition, blood parameters and performance of lactating goats and their kids were evaluated.

MATERIALS AND METHODS

The field experiments were carried out at Goat Research Unit of the Department of Animal Production at El-Noubaria Experimental Farm, National Research Centre, Noubaria, Egypt.

Five experimental diets were formulated as follow:

T1: Concentrate feed mixture (CFM) + berseem hay (BH) as control diet,

T2: CFM + untreated SBP + BH, T3: CFM + SBP treated with 2% urea on DM basis+ BH, T4: CFM + SBP treated with 4% urea on DM basis + BH and T5: CFM + SBP treated with 6% urea on DM basis+ BH.

The concentrate to roughage ratio was 70: 30%; the ratio of SBP was 30% of CFM.

The experiment lasted for 140 days (one month before kidding and three months as lactation and weaning period). Forty female Barki goat does (about 4 years old and weighing 41±1.5 kg) at the fifth month of pregnancy were randomly divided into five groups of 8 heads in each. Does were fed on the experimental diets for a month before kidding, and after kidding goats were weighed as the initial body weight (31±1.7 kg) and fed in the same diets till the end of lactation stage. Goat does were fed their daily diet according to average body weight, which was adjusted continuously every two weeks according to Kearl (1982). The concentrate and roughage were offered twice daily at 7 a.m. and 1 p.m. The offered and the refusals were weighed daily and the goats were weighed every two weeks. Fresh water had excess to the goats twice daily at 7 a.m. and 1 p.m.

At the end of lactation stage, a digestibility trial was carried out to study the effect of feeding the experimental rations on digestibility coefficients, rumen parameters, microbial protein, protozoal count, and some blood costituents. Four does from each group were placed in metabolic cages, weighed at the start and the end of the trial. The trial lasted for 20 days from which the first 15 days were considered as an adaptation and preliminary

period, followed by 5 days as collection period. Over the collection period, daily amount of feed consumed, residuals, feces, urine and drinking water were estimated for each doe, and samples of ruminal liquor and blood were taken.

Urea treatment:

Air-dried SBP was moistened for 40% DM and sprayed with urea solution at levels of 2, 4 and 6% plus 10% molasses from the dry matter then it mixed well and placed in plastic light bags (1.2 m x 0.8 m and 0.4 mm thickness) with capacity of about 50 kg. Each bag was pressed well, covered well by heavy stones and kept closed for 21 days. Thereafter, the bags were shuffled upside-down and mixed well every day to dry in the sun and become acceptable for animals.

Proximate analysis:

The proximate analysis of ration samples was carried out according to the A.O.A.C. (1990) to determine contents of DM, CP, CF and EE. While, NFE content was obtained by the difference. However, NDF, ADF and ADL contents were determined according to the procedures of Van Soest *et al.* (1991). Cellulose and hemicelluloses were calculated by the difference between NDF and ADF for hemicelluloses, ADF and ADL for cellulose.

Rumen liquor parameters:

Rumen liquor samples were obtained at 0, 3 and 6 hours postfeeding. Ruminal pH value was immediately measured with pH meter, while concentrations of ammonia nitrogen, total nitrogen and non-protein nitrogen were determined by the modified semi-micro-kjeldahl digestion method according to A.O.A.C (1990). Concentration of true protein nitrogen was calculated by subtracting. Concentration of total volatile fatty acids (TVFAs) was determined according to Warner (1964). Ruminal microbial protein was estimated as described by Makkar *et al.* (1982).

For classification and determination of ruminal ciliated protozoal count, the filtered rumen liquor were collected at 0, 3 and 6 hours post feeding and stained with 4 times volume of methyl-green formalin saline solution as described by Ogimoto and Imai (1981), then stoked in dark place until examination. After gentle mixing of fixed rumen liquor samples, one drop was poured on hemocytometer slide, covered with a cover slip and examined under a light microscope for identification of genera and species of microorganisms according to the description published by Dehority (1993). **Sampling of blood:**

Goats from each treatment were used to obtain 12 ml blood from the jugular vein at zero and 4 hours post-feeding. Blood samples of lambs were left to coagulate at room temperature, then centrifuged at 4000 rpm for 15 min to separate serum and kept it frozen at -20°C till analyses of concentrations of total proteins (determined by using electronic apparatus), albumin (Doumas and Biggs,1971), globulin (was obtained by subtracting) and urea (Patton and Crouch, 1977). Activity of aspartate aminotransferase (AST) and alanin aminotransferase (ALT) was determined according to Reitman and Frankel (1957).

Sampling and analysis of milk:

Goats were milked twice a day at 8.00 a.m. and 5.00 p.m. during the last three days of each month of lactation period. Milk samples were immediately collected from each animal after morning and evening milkings and milk yield was recorded. The sample of each animal represented a mixed sample of constant percentage of the morning and evening yield. Milk samples were analyzed for total solids (TS), fat, total protein (TP), lactose and ash by Bentley 150 infrared milk analyzer (Bentley Instruments, Chaska, MN, USA) according to A.O.A.C. (1990) procedures. Solids-not-fat (SNF) was calculated by subtracting fat from total solids percentage. Fat corrected milk (4% fat) was calculated by using the following equation according to Gaines (1928):

FCM = 0.4 M + 15 F, Where: M = milk yield (g) and F = fat yield (g).

Statistical analysis:

Data was statistically analyzed according to statistical analysis system of (SAS, 2000). Data of digestibility coefficients, nitrogen balance and water balance were analyzed by one-way analysis and the model was:

 $Y_{ij} = M + T_i + e_{ij}$

The used design for rumen fermentations, blood samples, body weight, feed intake and milk yield and composition was two-way analysis and the following model was used:

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

Where: Y_{ij} = experimental observation, μ = general mean, T_i = effect of experimental diet (i =1:5), I_j = effect of sampling time (j=0, 3 and 6 or 0 and 4), T_{lij} = effect of interaction of experimental diet and sampling time, and e_{ij} = experimental error.

Separation among means was carried out by using Duncan's multiple test (Duncan, 1955). The overall level for statistical significance was set at $P \le 0.05$. All values were expressed as statistical means \pm standard error of means (SEM)

RESULTS AND DISCUSSION

Chemical composition and cell wall constituents:

Data represented in Table (1) showed that SBP treated with urea (T3, T4 and T5) enhanced chemical composition and cell wall constituents more than untreated SBP and control. Experimental treatments including T5, T4 and T3 increased DM, EE, NFE and CP contents more than untreated diet (T2). While, the same diets had less Ash, CF, NDF, ADF, ADL, cellulose and hemicellulose contents as compared to untreated diet, except for T4 and T2, which had almost similar contents of cellulose. In addition, urea treatments increased DM, Ash and CP contents as compared to control diet (T1), although, T1 had higher values of OM, EE, CF and NFE more than urea treated diets and untreated diet (T2). Also, T1 had less NDF, ADF, ADL, cellulose and hemicellulose contents as compared to treated and untreated diets.

It is clear that SBP treated with 4 and 6 % urea (T4 and T5) had almost similar chemical composition and cell wall constituents, but treated diets with 4 and 6% urea (T4 and T5, respectively) were better than that treated with 2% urea (T3) in all nutrient percentage.

Results of untreated SBP are in agreement with the results reported by Saleh *et al.* (2001), Talha *et al.* (2002), Abedo *et al.* (2005) and Aziz (2014). Regarding the chemical composition, the increased CF content in untreated diet through partial replacement of CFM by SBP could be attributed to the high content of CF in SBP (17–22%) (Papadomichelakis *et al.* 2004).

Similar results were obtained by Singh *et al.* (1990), who treated wheat straw by 4% urea solution and reported a decreases in CF, NDF, ADF, hemicellulose and lignin contents in urea treated wheat straw. Mohamed (1998) and Koening and Beauchemin (2005) reported that urea supplementation in diets increased CP content and digestibility. Also, Aziz (2009) found that olive tree byproducts treated with 4% urea increased DM, OM, EE and CP contents and decreased CF, NDF, ADF, DAL, cellulose and hemicellulose contents as compared to control group in sheep. Elkholy *et al.* (2009) observed an elevation in CP and reduction in CF, ADF, and NDF contents in corn crop residues treated with 4% urea. Moreover, Okab *et al.* (2012) reported that OM, EE, CF, NDF, ADF, DAL, cellulose and hemicellulose contents were improved in ration containing 50% CFM+50% SPB treated with 4% urea as compared to untreated and control diets.

Table (1): Effect of urea treatments on chemical composition of the experimental diet.

(experimental diet.												
Item		Expe	erimenta	l diet		F	eedstu	ff					
item	T1	T2	Т3	T4	T5	CFM	Hay	USBP					
DM (%)	92.28	89.04	91.5	93.24	93.45	93.60	90.9	90.88					
Chemical	Chemical composition (%):												
OM 92.03 88.73 89.70 91.10 91.26 92.1 88.08 95.36													
Ash	7.97	11.27	10.3	8.90	8.74	7.90	11.92	4.64					
EE	3.15	2.16	2.55	2.78	2.79	3.12	2.45	1.30					
CP	12.20	10.87	13.40	17.00	17.16	12.42	13.78	9.18					
CF	14.39	17.91	13.74	12.60	12.48	11.56	24.90	24.64					
NFE	62.29	57.79	60.01	58.72	58.83	65.00	46.95	60.24					
Cell wall c	onstitu	ents (%)):										
NDF	30.98	47.54	43.63	41.54	41.47	31.14	63.29	60.39					
ADF	17.82	24.16	22.82	20.05	19.97	17.84	44.79	28.92					
ADL	4.87	4.15	3.88	3.20	3.13	5.00	7.27	2.78					
Cellulose	13.16	23.38	20.81	23.58	21.50	13.30	18.50	31.47					
Hemicell.*	12.95	20.01	18.94	16.85	16.84	12.84	37.52	26.14					
T1 (control): (oncontra	ato food m	ixturo (CE	M) + Bors	oom hav l	BH)							

T1 (control): Concentrate feed mixture (CFM) + Berseem hay (BH).

T2: CFM + untreated SBP + BH.

T3: CFM + SBP treated with 2% urea on DM basis+ BH.

T4: CFM + SBP treated with 4% urea on DM basis + BH.

T5: CFM + SBP treated with 6% urea on DM basis + BH.

USBP: Untreated SBP. Hemicell.* : Hemicellulose.

Goat performance during kidding:

Data in Table (2) indicated similarity in kidding rate without abortion or stillbirth cases in all groups. Also, number of born kids and twining cases without mortality cases were found after birth in goat kids of all groups.

Table (2): Effect of experimental diet on goat performance during kidding.

Item		Experimental diet								
Item	T1	T2	Т3	T4	T5					
Lambing rate (%)	100	100	100	100	100					
Number of kids	9	9	9	9	9					
Twining rate (%)	12.5	12.5	12.5	12.5	12.5					
Mortality	0	0	0	0	0					

Body weight and feed intake of does during lactation:

Data in Table (3) showed significant ($P \le 0.05$) effect of the experimental diet on live body weight of goat does, being higher ($P \le 0.05$)in goat group fed SBP treated with 2, 4 and 6% (T4 and T5) as compared to those fed untreated diet (T2) during early, mid and late lactation stages. In this respect, goats fed T4 and T5 showed significantly ($P \le 0.05$) the highest live body weight with insignificant differences between them. Overall mean of live body weight during all lactation stages was had the same trend Overall mean of live body weight increased ($P \le 0.05$) by progressed time of lactation stages in all groups. Data of body weight change during lactation stages indicated that urea treatments (T4 and T5) showed the highest ($P \le 0.05$) live body weight, followed by control group then untreated group, although the difference between T3 and control group was not significant ($P \le 0.05$), also the difference between control and untreated group was not significant.

The increment in body weight might be related to the reported efficient anaerobic rumen fermentation (Mohamed and Abou-Zeina 2008). This result was supported by the results of Aziz (2009), who demonstrated that the greatest body weight change was produced in sheep fed on olive tree byproducts treated with 4% urea compared to control group.

Data in Table (3) indicated significant ($P \le 0.05$) difference in feed intake as g/h/d, g/kg BW and g/kg BW^{0.75}), being higher in goats fed urea treated diets than untreated and control diet. At early lactation stage, T3 and T5 had significantly ($P \le 0.05$) increased feed intake (g/h/d) with no significant difference, followed by T4 and control group with no significant ($P \le 0.05$) difference, while T2 had the lowest ($P \le 0.05$) value. At mid lactation stage, T5 had the highest ($P \le 0.05$) value (g/h/d), while the difference among T4, control, T3 and T2 was not significant ($P \le 0.05$). At late lactation stage, T5 was the highest ($P \le 0.05$), followed by T4 and T3, then control and T2, although, the differences were insignificant among T5, T4 and T3 or among T4, T3, T1 and T2.

Overall mean of feed intake showed that T3 had the highest values (g/h/d, g/kg BW and g/ kg $BW^{0.75}$) followed by T5, T4, T1 then T2. Overall mean of feed intake at lactation stages indicated significant (P≤0.05) increase

by progressed stage of lactation. Increasing feed intake may be due to good palatability of urea treated SBP.

The present results were supported by Salman *et al.* (1998), who reported that feed intake significantly (P<0.05) increased by two urea ensiling methods (wet or dried) for corn stalks ration. Also, Galina *et al.* (2004) fed goat kids on corn stubble and alfalfa hay treated with 5% urea, they found that feed intake was increased in treated groups. Aziz (2009) indicated that feed intake was increased in sheep fed on olive tree by-products treated with 4% urea as compared to control group. Moreover, Okab *et al.* (2012) showed that daily feed intake was decreased (P<0.05) in lambs fed on the diet containing untreated SBP as compared to those fed on diet contained SBP treated with 4% urea and control diets. In this way, several studies found an increase in DM intake from sugar beet replacement (Olfaz *et al.* 2005; Omer *et al.* 2013).

Table (3): Effect of experimental diet on body weight and feed intake during lactation period.

Item		Ехр	erimenta	al diet		±SEN	Overall		
Item	T1	T2	Т3	T4	Т5	TOCM	mean		
		Live bo	dy weigh	nt (LBW,	kg):				
Early lactation	31.50 ^b	30.83 ^c	31.31 ^b	32.11 ^a	32.20 ^a		31.59 [°] ±0.051		
	32.27 ^b		32.33 ^b	33.31 ^a			32.59 ^b ±0.051		
Late lactation	33.53 ^b		33.50 ^b	35.15 ^a	35.12 ^a	0.116	34.00 ^a ±0.051		
	32.43 [⊳]	31.76 [°]	32.38 ^b	33.52 ^a	33.53 ^a	0.067	-		
Change in LBW	2.03 ^{bc}	1.90 ^c	2.21 ^b	3.01 ^a	3.02 ^a	0.096	-		
			ed intake	e (g/h/d):					
	293.9 ^b				1319.2 ^ª		296.0 ^c ±3.299		
	386.4 ^b		392.4 ^b		1457.2 ^ª	7.376	398.0 ^b ±3.299		
Late lactation	473.6 ^b	465.0 ^b	490.9 ^{ab}		1502.0 ^ª	7.376	482.3 ^ª ±3.299		
Overall mean	384.6 ^c	359.0 [°]	412.3 ^a	378.6 ^b	1426.1ª	.259	-		
			intake (g						
	41.09 ^b		43.22 ^ª		40.97 ^b		41.03 [°] ±0.112		
	42.97 ^c		43.06 ^b		43.80 ^ª		42.90 ^b ±0.112		
	43.96 ^c		44.50 ^b	42.11 ^e			43.63 ^a ±0.112		
Overall mean	42.67 ^b		43.59 ^ª	41.09 ^c		0.145	-		
Feed intake (g/ kg BW ^{0.75}):									
		94.65 ^c	102.12 ^a	94.70 ^c	97.61 ^b		97.39 ^c ±0.260		
			102.31 ^b		105.19 ⁴		102.53 ^b ±0.260		
Late lactation	106.44 ^t	107.06 ^a	106.67 ^b	102.57 ^d	104.11 ⁶	0.581	105.37 ^a ±0.260		
Overall mean	102.42 ^t	101.55 [♭]	103.70 ^a	98.83 ^c	102.30 ^t	0.335	-		

Means with different litters within each row or column are significantly different (P≤0.05).

Digestibility coefficients and nutritive values:

Data in Table (4) indicated that urea treatments (T4 and T5) significantly (P<0.05) increased body weight and feed intake of goats as compared to untreated and control groups, followed by T3 and T1 then T2, although the difference between T3 and control group was not significant

(P \leq 0.05) for body weight. Also, the differences among T4, T3, T1 and T2 were not significant (P \leq 0.05) for feed intake.

Data in Table (4) indicated significant ($P \le 0.05$) effect of urea treatment on digestibility coefficients of all nutrients and fiber fractions, being significantly (P < 0.05) higher for all urea treated diets (T3-T5) than control and USBP diets. However, USBP diet showed significantly (P < 0.05) lower digestibility coefficients and fiber fractions than the control diet. T4 and T5 had the highest values with no significant difference followed by T3, while the lowest digestibility coefficients were for untreated SBP then control group. There was no significant ($P \le 0.05$) difference among control and T2 in digestibility coefficient of CP and cellulose. This result indicated that rations containing untreated or ureated SBP had digestibility coefficients in rumen better or near to the digestibility coefficients of ration containing only concentrate feed mixture, this means that partial replacement of concentrate by SBP had good benefits.

Higher nutrients digestibility as a result of urea treatment may be related to the microbial activities which solubilizing carbohydrate esters of phenolic monomers in the cell wall (Khampa *et al.* 2009). The high water holding capacity of dry SBP due to the existence of pectic substances, methyl and carboxyl groups in its molecular structure might be the reason of its better digestion (El-Badawi and El-Kady, 2006). Hall *et al.* (1998) recorded that SBP has a relatively high content of soluble and insoluble NDF, but it can be considered an energy concentrate because both soluble and insoluble NDF are highly digestible. Moreover, Nianogo *et al.* (1997) observed an increase in DM intake and OM digestibility of urea treated rice straw; it is attributed to the chemical changes, which are related to the breaking of the lignin-polysaccharide links, which favors the release of digestible components.

Data in Table (4) showed significant (P≤0.05) effect of the experimental diet on nutritive values. Nutritive value as TDN (g/h/d) was higher in T5 than in other treatments, followed by T4 then control and T3, while the lowest (P≤0.05) value was for untreated group. Nutritive value as TDN (g/kg BW, g/kg BW^{0.75} and % of DMI) had no significant (P≤0.05) differences among T5, T4 and control group, T3 came in the second class, while T2 had the lowest (P≤0.05) values.

Also, urea treatments significantly (P≤0.05) increased the values of DCP (g/h/d, g/kg BW, g/kg BW^{0.75} and % of DMI) more than control and untreated SBP, T5 had the highest (P≤0.05) values of DCP (g/h/d, g/kg BW and g/kg BW^{0.75}) followed by T4 then T3, while, T2 had the lowest (P≤0.05) values followed by control (T1). It seems that T4 came in the first class for DCP % of DMI followed by T5.

Itom		Expe	erimental	diet		±SEM
ltem	T1	T2	Т3	T4	T5	
Number of animals	4	4	4	4	4	-
Live body weight (kg)	34.03 ^b	33.35 [°]	34.00 ^b	35.65 ^a	35.63 ^a	0.084
Fee intake (g/h/d)	1473.58 ^b	1465.02 ^b	1490.92 ^{ab}	1480.11 ^{ab}	1502.02 ^ª	7.376
			fficient (%			
DM	79.17 [°]	78.04 ^d	82.84 ^b	86.41 ^a	85.81 ^a	0.349
OM	80.05 ^c	78.97 ^d	83.13 ^b	87.03 ^a	86.79 ^a	0.193
EE	87.50 ^c	86.34 ^d	88.04 ^b	89.80 ^a	89.77 ^a	0.320
CP	79.32 ^c	79.07 ^c	80.95 ^b	84.73 ^a	84.69 ^a	0.230
CF	68.28 ^c	67.38 ^d	71.76 ^b	76.34 ^a	76.24 ^a	0.541
NFE	83.13 ^c	82.45 ^d	84.04 ^b	85.89 ^a	85.56 ^a	0.219
		Fiber frac				
NDF	74.03 ^c	71.86 ^d	77.64 ^b	79.92 ^a	79.13 ^a	0.343
ADF	62.27 ^c	60.78 ^d	66.81 ^b	70.42 ^a	69.66 ^a	0.333
ADL	56.58 ^c	53.76 ^d	61.89 ^b	65.75 ^a	65.36 ^a	0.265
Cellulose	67.97 ^c	67.09 ^c	71.33 ^b	75.46 ^a	75.13 ^a	0.346
Hemicellulose	78.36 ^c	77.10 ^d	81.34 ^b	85.32 ^a	85.13 ^a	0.329
		Nutritive v				
TDN g/h/d	1259.58 ^c	1124.72 ^e	1212.37 ^d	1263.30 ^b	1268.03 ^a	0.232
TDN g/kg BW	34.28 ^a	30.85 ^c	33.33 ^b	34.44 ^a	34.78 ^a	0.230
TDN g/kg BW ^{0.75}	84.40 ^a	75.81 ^c	81.87 ^b	84.76 ^a	85.47 ^a	0.425
TDN % of DMI	88.52 ^a	81.41 ^c	84.99 ^b	88.18 ^a	88.18 ^a	0.117
DCP g/h/d	127.38 ^d	106.20 ^e	141.60 ^c	192.41 ^b	194.51 ^a	0.024
DCP g/kg BW	3.46 ^d	2.91 ^e	3.89 ^c	5.24 ^b	5.33 ^a	0.026
DCP g/kg BW ^{0.75}	8.53 ^d	7.16 ^e	9.56 ^c	12.91 ^b	13.11 ^ª	0.047
DCP % of DMI	79.32 ^d	79.08 ^e	80.96 ^c	84.73 ^a	84.69 ^b	0.021

 Table (4): Effect of experimental diet on nutrient digestibility and nutritive value of the experimental rations.

Means with different litters within each row are significantly different (P≤0.05).

Based on the foregoing results, urea treatments especially T5 and T4 increased TDN values more than control and untreated group, while the three urea treatments increased DCP values. These results reflected the values of nutrient digestibility for rations which indicated that urea treatments were more efficient in digestibility of nutrients compared with control and untreated SBP as that the improvement in nutritive values are associated with the increased digestion of fibrous materials and bacterial digestion of cell wall content (Hassan *et al.* 2005).

Similar results were obtained by Aziz (2009), who demonstrated an increase in digestibility coefficients of DM, OM, EE, CP, CF, NDF, ADF, DAL, cellulose and hemicellulose and an increase in TDN and DCP intake in sheep fed on olive tree byproducts treated with 4% urea compared to control and untreated group. Also, Salman *et al.* (2011) showed that 3% urea treatment of sugarcane had increased DM, CP, EE, and ash, while, decreased OM, CF and NFE.

Moreover, Okab et al. (2012) found that digestibility coefficients of CP, CF, and EE were decreased (P<0.05) in untreated compared to control diet. Although, the inclusion of treated SBP with 4% urea in the diet of sheep increased the digestibility coefficients of DM, OM and CP in comparison with control diet. However, CF digestibility coefficient was reduced (P<0.05) as compared to control diet. Also, they found that treated SBP diets increased (P<0.05) TDN and DCP intake as compared to untreated and control diets. On the other hand, El-Badawi and El-Kady (2006) noted that digestibility of CP and EE were decreased, while, CF digestibility was extremely higher in sheep fed ration containing 50% SBP sprayed with urea solution 30 g/kg SBP to compare traditional feed mixture. Nutritive value expressed in terms of TDN was higher by about 5% for the 50% ureated SBP more than control. Also, Omer et al. (2013) showed that inclusion of SBP in sheep ration insignificantly (P<0.05) improved DM and OM digestibilities, while, it significantly (P<0.05) increased CF digestibility, but, it significantly (P<0.05) decreased EE digestibility. Dietary treatment had no effect on nutritive values (TDN and DCP).

Nitrogen balance:

Data in Table (5) showed that urea treatments (T3, T4 and T5) increased (P \leq 0.05) nitrogen intake values (g/h/d, g/kg BW and g/kg BW^{0.75}) more than control and USBP groups. T5 had the highest (P \leq 0.05) values, followed by T4 then T3, while the lowest (P \leq 0.05) values were for untreated group (T2) then and control.

Urea treatments significantly (P≤0.05) increased fecal nitrogen excretion, T5 had the highest (P≤0.05) values of fecal nitrogen excretion (g/h/d, g/kg BW^{0.75} and % of NI), followed by T4 then T3, while the lowest (P≤0.05) value was for untreated group (T2). No significant difference was found among T5, T4, T3 and T1 in values of fecal nitrogen excretion (g/kg BW).

Urinary nitrogen excretion was differed from fecal N, untreated and control groups increased (P≤0.05) urinary N (g/h/d, g/kg BW, g/kg BW^{0.75} and % of NI) more than urea treatments, although, the difference between T1 and T2 except for values of urinary N % of N intake (NI). T4 had the lowest (P≤0.05) values followed by T5 then T3, although the difference between T4 and T5 was not significant (P≤0.05). Total nitrogen excretion values were the same trend of urinary N as that untreated and control groups (T2 and T1) had the highest (P≤0.05) values of (g/h/d, g/kg BW, g/kg BW^{0.75} and % of NI) with no significant (P≤0.05) except for values of total N excretion % of NI, while urea treatments especially T4 had the lowest values followed by T5 then T3, although the difference among the three treatments was not significant (P≤0.05).

It is clear that urea treatments decreased total N excretion, but, it increased (P≤0.05) nitrogen balance (g/h/d, g/kg BW, g/kg BW^{0.75},% of NI and % of DN) more than untreated and control groups, T4 had the highest (P≤0.05) values followed by T5 then T3, although the difference between T4 and T5 was not significant (P≤0.05), while untreated group(T2) had the lowest (P≤0.05) nitrogen balance values followed by control group (T1). It seems that urea treatments for SBP increased (P≤0.05) nitrogen balance by 12.74- 22.03

% of N intake more than untreated SBP, and by 6.62- 16.18 % of N intake more than control.

The superiority in N balance by feeding urea treated SBP might be due to the higher improvement of its OM and CP digestibility or due to the higher utilization of urea N and thus lower total nitrogen excretion or might be related to higher improvement in rumen fermentation.

Similar results were obtained by Mohamed *et al.* (1998) who indicated that N-balance of ureated rice straw (4% urea) had significantly (P<0.05) the highest value than untreated and ammoniated rice straw. El-Badawi *et al.* (2003) found that dietary nitrogen utilization (%N-balance of N-intake) was obviously higher (P<0.05) with feed mixtures containing 50 and 75% SBP. Aziz (2009) demonstrated that the greatest nitrogen balance was produced in sheep fed on olive tree by-products treated with 4% urea compared to control group. Also, Okab *et al.* (2012) reported that total N excretion was increased (P<0.05) in untreated SBP and control diets, while, N balance was greater (P<0.05) in diets containing 50% SBP treated with 4% urea compared to control and untreated diets. Moreover, Omer *et al.* (2013) indicated that N balance expressed as relative to N-Intake or digestible-N was significantly P<0.05) improved with lower (P<0.05) urinary N loss for sheep fed SBP feed mixture.

Palanaa	ltom		Expe	erimental	diet		±SEM
Balance	ltem	T1	T2	Т3	T4	T5	TOEIN
Nitrogon	g/h/d	25.70 ^d	21.49 ^e	27.98 ^c	36.33 ^b	36.75 ^a	0.005
Nitrogen intake	g/kg BW	0.700 ^d	0.592 ^e	0.767 ^c	0.992 ^b	1.010 ^a	0.005
IIIIake	g/ kg BW ^{0.75}	1.72 ^d	1.44 ^e	1.89 ^c	2.44 ^b	2.47 ^a	0.009
	g/h/d	5.32 ^d	4.50 ^e	5.33 ^c	5.55 [⊳]	5.62 ^a	0.001
Fecal	g/kg BW	0.145 ^ª	0.120 ^b	0.147 ^a	0.150 ^ª	0.150 ^a	0.002
nitrogen	g/ kg BW ^{0.75}	0.355 ^d	0.302 ^e	0.362 ^c	0.370 ^b	0.380 ^a	0.002
	% of N intake	20.68 ^d	20.92 ^e	19.04 ^c	15.27 ^b	15.31 ^a	0.003
	g/h/d	6.39 ^a	6.55 ^a	5.56 ^b	5.12 ^c	5.26 ^c	0.090
Urinary	g/kg BW	0.175 ^a	0.180 ^a	0.152 ^b	0.140 ^c	0.142 ^c	0.003
nitrogen	g/ kg BW ^{0.75}	0.427 ^ª	0.440 ^a	0.375 ^b	0.345 [°]	0.355 ^{bc}	0.007
	% of N intake	24.86 ^b	30.48 ^a	19.89 ^c	14.11 ^d	14.31 ^d	0.311
	g/h/d	11.71 ^a	11.05 ^b	10.89 ^{bc}	10.67 ^c	10.88 ^{bc}	0.090
Total N	g/kg BW	0.320 ^a	0.305 ^b	0.300 ^{bc}	0.290 ^c	0.300 ^{bc}	0.003
excretion	g/ kg BW ^{0.75}	0.782 ^ª	0.745 ^b	0.735 ^{bc}	0.717 ^c	0.735 ^{bc}	0.007
	% of N intake	45.55 ^b	51.40 ^a	38.93 ^c	29.38 ^d	29.61 ^d	0.311
	g/h/d	13.99 ^c	10.44 ^d	17.09 ^b	25.86 ^a	25.66 ^a	0.090
Nitrogen	g/kg BW	0.380 ^c	0.287 ^d	0.470 ^b	0.710 ^a	0.700 ^ª	0.004
balance	g/ kg BW ^{0.75}	0.935 ^d	0.705 ^e	1.155 [°]	1.745 ^ª	1.722 ^b	0.006
	% of N intake	54.44 ^c	48.59 ^d	61.06 ^b	70.62 ^a	70.38 ^a	0.311

Table (5): Nitrogen	balance	for	goat	groups	affected	by	experimental
diet.							

Means with different litters within each row are significantly different (P≤0.05). Water balance:

Data of Table (6) showed significant (P \leq 0.05) difference in free drinking water and total water intake (ml/h/d or ml/kg W^{0.82}). Control and

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untreated groups (T1 and T2) increased (P≤0.05) free drinking water more than urea treatments (T3, T4 and T5), although the differences between T1 and T2or among urea treatments were not significant. T4 had the lowest (P≤0.05) values, followed by T5 then T3. Untreated SBP group (T2) had the highest (P≤0.05) value of combined water (ml/h/d or ml/kg $W^{0.82}$), followed by T3, T1 then T5, while the lowest (P≤0.05) value was for T4. Metabolic water (ml/h/d or ml/kg $W^{0.82}$) was significantly (P≤0.05) differed, T5 had the highest (P≤0.05) metabolic water (ml/h/d), followed by T5, T1, T3 and T4, while the lowest (P≤0.05) value was for untreated SBP.

Itom		Experimental diet							
iten	1	T1	T2	Т3	T4	T5	Μ		
Free	ml/h/d	3925.0 ^a		3725.0 ^{ab}			67.27		
	ml/kg W ^{0.82}	263.2 ^a	262.9 ^a			187.6 ^c	5.092		
Combined	ml/h/d	109.3 ^c	151.4 ^a		96.8 ^d	93.8 ^e	0.148		
	ml/kg W ^{0.82}	7.32 ^c	10.20 ^a		6.50 ^d	-	0.040		
Metabolic	ml/h/d	869.1 ^c	776.1 ^e		871.7 [⊳]		0.300		
	ml/kg W ^{0.82}		52.3 ^c				0.301		
Total	ml/h/d	4903.4 ^a	4827.5 ^{ab}		4553.5 [°]	4578.7 [°]	67.30		
	ml/kg W ^{0.82}								
Urinary	ml/h/d								
water					30.78 ^{bc}				
	% of intake	11.80 ^{ab}	12.22 ^a		10.40 ^{ab}				
Fecal	ml/h/d	87.63 ^a	88.24 ^a		82.02 ^c		0.464		
water:			5.95 ^a	5.66 ^b	5.50 ^c	4.30 ^ª	0.046		
	% of intake	1.79	1.82	1.79	1.80		0.022		
Total	ml/h/d	666.38 ^a		596.31 ^{ab}	540.77 ^b				
water				40.26 ^{ab}	36.29 ^{bc}				
execration	% of intake	13.59 ^b	14.05 ^a	12.74 [°]	11.87 ^d	12.15 ^{cd}	0.694		
	ml/h/d	4237.00							
er balance				275.94	269.26				
		00.40			88.12 ^a		0.694		
	Free Combined Metabolic Total Urinary water Fecal water: Total water execration er balance	ml/kg W ^{0.82} Combined ml/h/d ml/kg W ^{0.82} Metabolic ml/h/d ml/kg W ^{0.82} Total ml/h/d ml/kg W ^{0.82} Urinary ml/h/d ml/kg W ^{0.82} Vrinary ml/h/d ml/kg W ^{0.82} Voinary ml/h/d ml/kg W ^{0.82} Voinary ml/h/d ml/kg W ^{0.82} Yo of intake ml/h/d ml/kg W ^{0.82} Total ml/h/d ml/kg W ^{0.82} Yo of intake ml/h/d ml/kg W ^{0.82}	Free ml/h/d ml/kg W ^{0.81} 3925.0 ^a 263.2 ^a Combined ml/h/d ml/kg W ^{0.82} 109.3 ^c 7.32 ^c Metabolic ml/h/d ml/kg W ^{0.82} 869.1 ^c 58.2 ^a Total ml/h/d ml/kg W ^{0.82} 869.1 ^c 58.2 ^a Urinary ml/h/d ml/kg W ^{0.82} 578.8 ^{ab} 38.84 ^a 11.80 ^{ab} Fecal ml/h/d ml/kg W ^{0.82} % of intake 57.8.7 ^a 1.79 Total ml/h/d ml/kg W ^{0.82} % of intake 5.87 ^a 1.79 Total ml/h/d ml/kg W ^{0.82} % of intake 666.38 ^a 44.72 ^a 13.59 ^b execration ml/h/d ml/kg W ^{0.82} % of intake 4237.00 284.00 86.40 ^b	Item T1 T2 Free ml/h/d ml/kg W ^{0.8} 3925.0 ^a 3900.0 ^a Combined ml/h/d ml/kg W ^{0.82} 263.2 ^a 262.9 ^a Combined ml/h/d ml/kg W ^{0.82} 109.3 ^c 151.4 ^a Metabolic ml/h/d ml/kg W ^{0.82} 869.1 ^c 776.1 ^e Metabolic ml/h/d ml/kg W ^{0.82} 869.1 ^c 776.1 ^e Total ml/h/d ml/kg W ^{0.82} 4903.4 ^a 4827.5 ^{ab} Vinary ml/h/d ml/kg W ^{0.82} 578.8 ^{ab} 590.0 ^a water ml/h/d ml/kg W ^{0.82} 87.63 ^a 88.24 ^a Total ml/h/d ml/kg W ^{0.82} 57.63 ^a 88.24 ^a water: ml/h/d % of intake 87.63 ^a 88.24 ^a Total ml/h/d ml/kg W ^{0.82} 663.8 ^a 678.24 ^a water ml/kg W ^{0.82} 44.72 ^a 45.73 ^a execration ml/h/d % of intake 284.00 279.69 water ml/kg W ^{0.82} 284.00 279.69 % of intake 86.40 ^b 85.95 ^c	$\begin{array}{ c c c c c c c c c c c c c c c c c c c$	T1 T2 T3 T4 Free ml/h/d ml/kg W ^{0.8} 3925.0 ^a 3900.0 ^a 3725.0 ^{ab} 3585.0 ^b Combined ml/h/d ml/kg W ^{0.82} 263.2 ^a 262.9 ^a 251.5 ^{ab} 240.6 ^b Combined ml/h/d ml/kg W ^{0.82} 7.32 ^c 10.20 ^a 8.18 ^b 6.50 ^d Metabolic ml/h/d ml/kg W ^{0.82} 869.1 ^c 776.1 ^e 836.5 ^d 871.7 ^b Total ml/h/d ml/kg W ^{0.82} 4903.4 ^a 4827.5 ^{ab} 4682.8 ^{bc} 4553.5 ^c Total ml/h/d ml/kg W ^{0.82} 328.7 ^a 325.4 ^a 316.2 ^{ab} 305.6 ^b Urinary ml/h/d 578.8 ^{ab} 590.0 ^a 512.5 ^{abc} 458.8 ^c water ml/h/d 87.63 ^a 88.24 ^a 39.78 ^a 34.60 ^{ab} 30.78 ^{bc} water: ml/h/d 87.63 ^a 88.24 ^a 596.31 ^{ab} 540.77 ^b water ml/h/d 87.63 ^a 88.24 ^a 596.31 ^{ab} 540.77 ^b water ml/h/d 87.63 ^a 678.24 ^a	$\begin{array}{ c c c c c c c c c c c c c c c c c c c$		

 Table (6): Water balance for goat groups affected by experimental diet.

Means with different litters within each row or column are significantly different (P≤0.05).

It was found that urinary, fecal and total water execration showed similar trends, untreated group (T2) had the highest (P≤0.05) values, followed by control group (T1), while all urea treatments had the lowest (P≤0.05) values, although the difference among urea treatments was not significant (P≤0.05), T4 had the lowest values followed by T5 then T3. Water balance (ml/h/d and ml/kg $W^{0.82}$) showed insignificant (P≤0.05) differences among treatments, although the data of water balance (% of intake) showed significant (P≤0.05) difference; urea treatments had higher water balance than control and untreated groups, although the differences among urea treatments were not significant (P≤0.05). T4 had the highest (P≤0.05) water balance, followed by T5 then T3, while untreated group (T2) had the lowest water balance followed by control group (T1).

In agreement with the present results, Deyab (1995) and Hassona (1997) showed that the total daily water consumption and water balance was significantly higher in animals fed ureated rice straw than untreated rice straw (control). Also, Aziz (2009) demonstrated that total water intake and water balance (ml/h/d or ml/kg $W^{0.82}$) were higher in sheep fed on olive tree by-products treated with 4% urea.

Rumen parameters:

Ruminal pH value, volatile fatty acids (VFAs) and molar proportion of individual VFAs:

Data in Table (7) showed that urea treatments decreased (P \leq 0.05) ruminal pH values as compared to control and untreated groups; the lowest (P \leq 0.05) ruminal pH values were for T4, followed by T5 with no significant difference (P \leq 0.05) then T3. While, control group (T1) had the highest (P \leq 0.05) ruminal pH value, followed by untreated group (T2).

Overall means of ruminal pH at different sampling times showed the highest value at zero time (per-feeding), and then it showed a significant decrease ($P \le 0.05$) 3 h post-feeding, then increased with progressed time of feeding 6 h post-feeding.

Urea treatments significantly ($P \le 0.05$) increased total VFAs concentration as compared to untreated and control groups. The highest value was for T4, followed by T5 then T3, although the differences between T4 and T5 or between T5 and T3 were not significant, while the lowest value was for untreated group and control group with no significant difference between them. It is clear that ruminal VFAs concentration increased when ruminal pH value was decreased at different sampling times. This trend may be due to ruminal fermentation by rumen microorganisms (Dawson *et al.* 1990).

A significant differences were detected in molar proportions of ruminal individual VFAs. Urea treatments significantly increased (P \leq 0.05) molar percentage of acetic, propionic and butyric more than control and untreated group. The highest (P \leq 0.05) values were for T4, followed by T5 then, although the difference between T4 and T5 was not significant for propionic value. While, the lowest values were for untreated group (T2), followed by control group (T1). Untreated group, followed by T3 then control group significantly (P \leq 0.05) had higher values of acetic to propionic ratio more than T4 and T5, being the highest (P \leq 0.05) in T2, while, T4 had the lowest (P \leq 0.05) value. The differences among T2, T3 and T1 or between T4 and T5 were not significant.

Overall means of total VFAs concentration and molar proportions of acetic, propionic and butyric at different sampling times clearly showed that the lowest values were at zero h (per-feeding) then showed a significant increase ($P \le 0.05$) to reach the highest value 3 h post-feeding then it showed decrease with progressed time of feeding 6 h post-feeding. While, overall mean of A/P at different sampling times was not significant. The increase in TVFAs 3 h post-feeding might be related to the more utilization of dietary energy and positive fermentation in the rumen (Tagari *et al.* 1964). Also, increasing ruminal TVFAs concentration is an indicator for better utilization of dietary carbohydrates (Fadel *et al.* 1987).

ltem	Time		-	rimenta			±SEM	Overall
item	(h)	T1	T2	Т3	T4	T5		mean
Ruminal	0	7.30	6.77	6.67	6.42	6.42	0.099	
pH value	3	6.32	5.95	5.87	5.77	5.87	0.099	
pri value	6	6.52	6.37	6.17	5.90	5.97	0.099	6.19 ^c ±0.044
Overall mea	an	6.71 ^a	6.36 ^D	6.24 ^{bc}	6.03 ^ª	6.09 ^ª	0.057	
TVFAs(ml	0	7.93	7.84	8.21	8.56	8.41	0.139	8.19 ^c ±0.062
equiv/100	3	9.75	9.57	11.59	12.14	11.93	0.139	10.99 ^ª ±0.062
ml R.L)	6	8.97	8.56	10.01	10.18	10.11	0.139	9.57 ^b ±0.062
Overall mea	an	8.88 ^c	8.65 ^c	9.94 ^b	10.29 ^a	10.15 ^{ab}	0.080	-
	Мо	olar pro	portior	ns of in	dividua	al VFAs		
	0	33.20	32.41	35.95	38.44	37.62		35.52 ^c ±0.085
Acetic	3	38.88	37.80	41.56	42.68	42.39		40.66 ^ª ±0.085
	6	35.87	35.74	38.89	39.94	39.54	0.192	38.00 ^b ±0.085
Overall mea	an	35.99 ^d	35.32 ^e	38.80 ^c	40.35 ^a	39.85 ^b	0.110	-
Propionic	0	17.26	16.94	18.87	21.25	20.48		18.96 ^c ±0.300
FIOPIOIIIC	3	21.47	19.18	20.55	26.56	26.41		22.83 ^ª ±0.300
	6	19.27	17.75	21.74	23.79	23.54	0.672	21.22 ^b ±0.300
Overall mea	an	19.33 ^b	17.95 [°]	20.39 ^b	23.87 ^a	23.48 ^a	0.388	-
Putyrio	0	15.29	15.22	16.38	18.62	18.19		16.74 [°] ±0.091
Butyric	3	17.75	17.54	18.82	22.65	22.12		19.77 ^ª ±0.091
	6	16.45	15.50	18.92	19.58	19.42	0.205	17.97 ^b ±0.091
Overall mea	an	16.49 ^d	16.09 ^e	18.04 ^c	20.28 ^a	19.91 ^b	0.118	
A/D ratio	0	1.92	1.91	1.90	1.80	1.83	0.091	1.87±0.040
A/P ratio	3	1.81	1.97	2.15	1.60	1.60	0.091	1.82±0.040
	6	1.86	2.01	1.78	1.68	1.68	0.091	1.80±0.040
Dverall mean 1.86 ^a 1.96 ^a 1.94 ^a 1.69 ^b 1.70 ^b 0.052 -								
leans with dif	fferent lit	ters with	in each r	ow or co	olumn are	e signific	antly diff	erent (P≤0.05).

 Table (7): Effect of experimental diet on ruminal pH, volatile fatty acids and molar proportion of individual VFA's.

The present results are in agreement with those obtained by Fouad *et al.* (1998), who showed that ruminal pH values of control group was significantly (P<0.05) higher than urea treatment. Also, Bodas *et al.* (2007) found that inclusion of SBP in cereal based diets for fattening lambs seems to enhance the ruminal environment and prevent ruminal acidosis. Also they reported that ruminal pH was significantly (P<0.05) lower in lambs fed on the control concentrate compared to concentrate with SBP. A significant decrease (P<0.05) in total VFAs concentration occurred when SBP was included in the concentrate. Omer *et al.* (2013) indicated that inclusion untreated SBP in sheep ration significantly (P<0.05) increased total VFAs concentration, however it had no significant effect on pH value compared to control ration (CFM).

Total nitrogen, true protein nitrogen, non-protein nitrogen, ammonia nitrogen and microbial protein concentrations:

Urea treatments significantly (P≤0.05) increased total nitrogen, true protein, non-protein nitrogen, ammonia nitrogen and microbial protein

concentrations more than control and untreated group (Table 8). T4 had the highest (P \leq 0.05) values of total nitrogen, true protein, non-protein nitrogen (NPN) and microbial protein concentrations, followed by T5 then T3, while, the highest (P \leq 0.05) value of ammonia concentration was for T5, followed by T4 then T3. Untreated group (T2) had the lowest (P \leq 0.05) values of all parameters, followed by control group (T1).

Data showed insignificant differences among T4, T5 and T3 for total nitrogen value. Also, there was insignificant difference between T4 and T5 for true protein and NPN values. Overall means of TN, TP, NPN, NH3-N and microbial protein concentrations at different sampling times showed that the lowest ($P \le 0.05$) values were at zero h (per-feeding) then significantly ($P \le 0.05$) increased to reach the highest ($P \le 0.05$) values 3 h post-feeding, then it decreased ($P \le 0.05$) with progressed time of feeding 6 h post-feeding.

It is clear that the level of 4% urea (T4) enhanced all rumen parameters concentrations more than level of 6% urea (T5), except that for ammonia nitrogen concentration, it was higher in T5. This is may be due to the highest digestibility of CP and the highest nitrogen balance in T4 more than T5. Also, higher microbial protein concentration in T4 indicated that level of 4% urea was more efficient.

The improvement in rumen parameters by urea treatments may be due to the increase in digestibility coefficients of all nutrients, or may be related to the improvement in microbial population in the rumen which plays an important role in rumen fermentation. The reduction of ammonia nitrogen in the rumen liquor by progressed time of feeding appears to be the result of increased incorporation of ammonia nitrogen into microbial protein and it was considered as a direct result to stimulated microbial activity (Tagari *et al.* 1964).

Silva and Ørskov (1988) reported that feeding SBP might encourage growth of cellulolytic and hemicellulolytic microorganisms and this, in turn, should increase the extent of forage digestion. On the other hand, there was a good efficiency in ruminal protein synthesis when a non protein nitrogen source was in feeds containing rapidly fermentable carbohydrates (Lanza *et al.* 2001). Therefore, use of SBP may have increased ruminal protein synthesis.

The present results are supported by the results of Mousa (2003) fed Baladi male lambs with untreated or ureated barley straw (4% urea) he found that NH₃-N concentrations in all groups increased 3 h post-feeding. Also, Gado *et al.* (2007) reported that the ruminal ammonia nitrogen levels were higher in sheep fed ureated yellow corn or barely grains at early hours after feeding than in those fed diets containing true protein (controls). Moreover, Aziz (2009) demonstrated a decrease in ruminal pH, and an increase in ruminal TVFAs, total nitrogen, true protein, non-protein nitrogen and ammonia nitrogen concentrations in sheep fed on olive tree by-products treated with 4% urea compared to control and untreated group. On the other hand, Omer *et al.* (2013) indicated that inclusion untreated SBP in sheep ration significantly (P<0.05) decreased ammonia nitrogen concentration. Bodas *et al.* (2007) found that no significant differences (P<0.05) were observed in ammonia-N

concentration when untreated SBP was inclusion in cereal based diets for fattening lambs.

Table (8): Effect of experimental diet on total nitrogen, true protein nitrogen, non-protein nitrogen, ammonia nitrogen and microbial protein concentrations (mg/100 ml RL).

microbial protein concentrations (ing/100 mi RL).										
Item	Time		Expe	rimenta	l diet		±SEM	Overall mean		
item	(h)	T1	T2	Т3	T4	T5	TOEIN	Overall mean		
Total	0	97.47	93.44	112.44	114.85	114.20	1.643	106.48 ^c ±0.734		
	3	122.81	117.66	131.37	134.20	133.51	1.643	127.91 ^ª ±0.734		
nitrogen	6	112.73	109.07	122.18	124.07	123.58	1.643	118.32 ^⁰ ±0.734		
Overall me	ean	111.00 ^b	106.72°	122.00 ⁸	124.37 ^a	123.76 ^a	0.948	-		
True	0	37.72	35.73	41.96	48.74	48.18	1.433	42.46 ^c ±0.640		
protein	3	44.72	40.63	47.94	53.62	52.84	1.433	47.95 ^ª ±0.640		
nitrogen	6	40.63	37.14	43.81	50.86	50.70	1.433	44.63 ^b ±0.640		
Overall me	ean	41.02 ^c	37.83 ^e	44.57 ^b	51.07 ^a	50.58 ^a	0.827	-		
	0	59.75	57.71	66.01	70.48	78.36	2.027	64.01 ^c ±0.906		
NPN	3	78.09	77.03	80.66	83.43	80.58	2.027	79.96 ^a ±0.906		
	6	72.10	71.93	72.87	78.36	73.20	2.027	73.69 ^b ±0.906		
Overall me	ean	69.98 ^c	68.89 ^ª	73.18 [⊳]	77.42 ^a	77.38 ^a	1.170	-		
Ammonia	0	30.89	30.48	33.61	37.38	38.90	0.442	34.25 ^c ±0.198		
nitrogen	3	36.56	34.58	38.16	43.95	45.20	0.442	39.69 ^ª ±0.198		
nitiogen	6	33.10	32.20	35.05	39.91	41.81	0.442	36.41 ^b ±0.198		
Overall me	ean	33.52 ^d	32.42 ^e	35.61 ^c	40.41 ^b	41.97 ^a	0.255	-		
Microbial	0	65.62	65.34	70.25	74.86	73.89	0.333	69.99 ^c ±0.149		
protein	3	109.16	108.81	113.38	121.22	120.11	0.333	114.54 ^ª ±0.149		
protein	6	87.47	86.60	93.67	98.04	97.94	0.333	92.74 ^b ±0.149		
Overall me	an	87.41 ^ª	86.92 ^c	92.43 ^c	98.04 ^a	97.31 [°]	0.192	-		

Means with different litters within each row or column are significantly different (P≤0.05).

Ruminal ciliated protozoa:

The identification of ruminal ciliate protozoa species and their density in the rumen liquor during different sampling times represented in Table (9). Data identified seven genera of ruminal protozoa in ruminal fluid of goats in this study. These genera (genus) are Entodinum spp., Epidinium ecaudatum, Diplodinum anisacanthum, Ophryoscolox spp., Polyolastron multivesiculatum, Isotrchia spp. and Dasytrachia rummantium. A significant increase (P≤0.05) was found in differential and total numbers of ruminal ciliated protozoa (x10⁴ cell/ml rumen liquor) with urea treatments more than control and untreated group, except for Diplodinum spp. which showed insignificant differences among all treatments. It is clear that T4 had the highest (P<0.05) density of all genera and total number of ciliated protozoa, followed by T5 then T3, while, the lowest (P≤0.05) values were for T2, followed by T1. The differences among T4, T5 and T3 for Epidinium spp., between T4 and T3 for Ophryoscolox spp. and between T5 and T3 for Polyolastron spp., Dasytrachia spp. and total count were not significant. Also, the differences between T1 and T2 for Epidinium spp., Ophryoscolox spp. and total count, and among all treatments in Isotrchia spp. were not significant. These results may be due to the effect of ruminal pH as that ruminal protozoa grow in a wide range of ruminal pH.

It is of interest to note that *Entodinum spp.* showed the highest number among all species. These results are in agreement with those reported by Ivan *et al.* (2000), who observed that *Entodinium* was the most detrimental of ciliated protozoa species.

Table	(9):	Effect	of	experimental	diet	on	count	of	ruminal	ciliated
		protozo	a (x	10 ⁴ cell /ml run	nen li	quo	r).			

protozoa (x10) cell /mi rumen liquor).											
ltem	Time			rimenta			±SEM	Overall			
item	(h)	T1	T2	Т3	T4	Т5		mean			
Entodinum	0	5.92	5.62	5.93	6.12	6.03	0.086	5.92 ^b ±0.038			
spp.	3	5.74	5.42	5.73	5.83	5.86	0.086	5.72 ^c ±0.038			
	6	6.24	6.02	6.70	8.00	7.05	0.086	6.80 ^a ±0.038			
Overall mea	an	5.96 ^d	5.69 ^e	6.12 ^c	6.65 ^a	6.32 ^b	0.049	-			
Enidinium	0	0.240	0.239	0.242	0.248	0.239	0.003	0.242 ^b ±0.001			
Epidinium	3	0.210	0.205	0.209	0.213	0.211	0.003	0.209 ^c ±0.001			
spp.	6	0.253	0.253	0.288	0.289	0.282	0.003	0.273 ^a ±0.001			
Overall mea	an	0.234 [°]	0.232 ^D	0.246 ^a	0.250 ^a	0.244 ^a	0.002	-			
Diplodinum	0	0.201	0.197	0.204	0.215	0.209	11.28	0.205 ^b ±5.047			
spp.	3	0.162	0.163	0.169	0.175	0.173	11.28				
	6	0.249	0.242	0.249	0.256	0.256	11.28	0.250 ^a ±5.047			
Overall mea	an	0.204	0.200	0.207	0.216	0.213	6.515	-			
Ophryoscolox	0	0.242	0.249	0.248	0.253	0.245	0.001	0.247 ^b ±0.008			
spp	3	0.209	0.206	0.211	0.210	0.208	0.001	0.209 ^c ±0.008			
	6	0.282	0.275	0.290	0.291	0.286	0.001	0.285 ^a ±0.008			
Overall mea	an	0.244 ^b	0.243 ^b	0.250 ^a	0.251 ^a	0.246 ^b	0.001	-			
Polyolastron	0	0.421	0.421	0.404	0.436	0.388	0.005	$0.414^{b} \pm 0.032$			
spp.	3	0.379	0.371	0.396	0.399	0.385	0.005	0.386 ^c ±0.032			
	6	0.408	0.420	0.459	0.482	0.469	0.005	$0.448^{a} \pm 0.032$			
Overall mea	an	0.403 ^c	0.404 ^c		0.439 ^a	0.414 ^b	0.003	-			
Isotrchia	0	0.275	0.279	0.275	0.283	0.270	800.0	0.276 ^b ±0.039			
spp.	3	0.243	0.244	0.238	0.243	0.236	800.0	0.241 ^c ±0.039			
	6	0.325	0.323	0.296	0.343	0.344	800.0	0.326 ^a ±0.039			
Overall mea	an	0.281 ^{ab}	0.282 ^{ab}	0.270 ^b	0.289 ^a	0.283 ^{ab}	0.005	-			
Desiduesta	0	0.449	0.484	0.503	0.510	0.500	0.007	$0.489^{b} \pm 0.035$			
Dasytrachia spp.	3	0.451	0.437	0.458	0.458	0.456	0.007	0.452 ^c ±0.035			
Spp.	6	0.538	0.544	0.551	0.620	0.552	0.007	0.561 ^a ±0.035			
Overall mea	an	0.479 ^c	0.488 ^c	0.504 ^b	0.529 ^a	0.503 ^b	0.004	-			
Total	0	7.75	7.50	7.80	8.07	7.89	11.27	7.80 ^b ±5.041			
protozoa	3	7.40	7.05	7.42	7.52	7.53	11.27	7.39 ^c ±5.041			
count	6	8.30	8.08	8.84	10.28	9.24	11.27	8.95 ^a ±5.041			
Overall mea	an	7.81 ^c	7.54 [°]	8.02 ^b	8.62 ^a	8.22 [⊳]	6.508	-			
loans with dif							41 11.00				

Means with different litters within each row or column are significantly different (P≤0.05).

In comparison among different sampling times, protozoal counts showed higher values per-feeding, then decreased (P \leq 0.05) 3 h post-feeding, then showed the highest (P \leq 0.05) numbers 6 h post-feeding. This trend was

in consistent with that of ruminal pH values, indicating that protozoal count is associated with ruminal pH value.

In the present study, the decreasing of protozoa number 3 h postfeeding may be related to the decreasing of ruminal pH after feeding as a result of increasing TVFAs concentration.

Similar results were obtained by Bhatia *et al.* (1992), who indicated that total protozoa count decreased 3 h after feeding and increased significantly 6 h post-feeding. Also, Aziz (2009) found that feeding sheep on olive tree by-products treated with 4% urea increased total ruminal protozoa count and the deferential number of *Entodinum, Isotrchia, Dasytrachia, Polyolastron, Epidinum,* and *Ophryoscolox spps.* as compared to control and untreated group. Moreover, Aziz (2014) reported that inclusion of biologically treated SBP in diets of sheep increased deferential and total ruminal protozoal count (x10⁴ cell /ml RL).

Blood composition:

A significant ($P\leq0.05$) increase was found in serum total proteins and albumin concentrations with urea treatments than control and untreated group. The highest ($P\leq0.05$) concentrations were recorded for T4, followed by T5 then T3, while, the lowest ($P\leq0.05$) values were recorded for untreated group (T2), followed by control group (T1). The differences between T5 and T3 for total proteins concentration, between T3 and T1 for albumin concentration were not significant. T3 showed the highest globulin concentration, followed by T1 then T2, while the lowest ($P\leq0.05$) concentrations were for T4 and T5. Albumin/globulin ratio increased in T1, T3 and T1 with no significant differences among them, being higher than in T4 and T5 with no significant differences between them (Table 10).

Serum urea concentration significantly ($P \le 0.05$) decreased with urea treatments more than control groups and untreated group, T4 had the lowest concentration, followed by T5 then T3, but the differences among the three treatments were not significant While, the highest concentration was for T2, followed by T1 with insignificant differences between them. The decrease in blood urea concentration in SBP groups might be due to partial reductions in provision of ruminal ammonia for subsequent hepatic conversion to urea (Mojtahedi and Mesgaran, 2011).

It was found that T4 only significantly ($P \le 0.05$) decreased activity of serum AST and ALT as compared to other treatments, which did not differ significantly. As affected by sampling time, all blood parameters 4 h postfeeding was higher ($P \le 0.05$) than pre-feeding values, except for albumin/globulin ratio which was not significant. These results showed that treating SBP with 2, 4 and 6% urea did not cause any lesions in liver and kidney functions of lactating goats.

The present results are supported by the results of Gado *et al.* (2007), who reported that concentration of total proteins, albumin, globulin and AL/GL ratio in sheep fed ureated yellow corn or ureated barely grains (4% urea) were higher comparing with control and untreated groups. Also, Aziz (2009) reported that serum total proteins, albumin, globulin concentrations were higher in sheep fed on olive tree byproducts treated with 4% urea compared to control and untreated group, while serum urea concentration, and activity of

AST and ALT decreased. On the other hand, Okab *et al.* (2012) found that hematological and biochemical parameters values of lambs fed on diets containing SBP treated with 4% urea were decreased (P<0.05) as compared to those fed on control and untreated diets. However, Abedo (2006), Bodas *et al.* (2007) and Mahjoubi *et al.* (2009) noticed insignificant differences between control and untreated SBP groups in any of the blood biochemical parameters studied. Also, Omer *et al.* (2013) and Sorathiya *et al.* (2015) reported that inclusion of untreated SBP in sheep diet significantly (P<0.05) decreased urea blood plasma concentration, however, it had no significant effect on other blood parameters compared to control.

activity in blood serum of goats.										
ltem	Time		Exper	imenta	l diet		±SEM	Overall		
item	(h)	T1	T2	T3	T4	T5	TOEM	mean		
Total proteins (g/dl)	0 4	8.07 8.90	7.86 8.28	8.69 9.33	8.64 10.12		0.109 0.109	8.40 ^b ±0.049 9.21 ^a ±0.049		
Overall mean		8.48 ^c	8.07 ^ª	9.01 [°]	9.38 ^a	9.08 ^D	0.077	-		
Albumin (g/dl)	0 4	4.40 4.68	3.88 4.53	4.42 5.09	5.39 6.30	5.42	0.105 0.105	4.75 ^b ±0.047 5.20 ^a ±0.047		
Overall mean	n	4.54 ^c	4.20 ^ª	4.75 ^c	5.85 ^a		0.074	-		
Globulin	0	3.67	3.98	4.27	3.25	3.06	0.119	3.64 ^b ±0.053		
(g/dl)	4	4.21	3.75	4.23	3.81		0.119	4.00 ^a ±0.053		
Overall mea	n	3.94 [⊳]	3.87 [¤]	4.25 ^a	3.53 [°]	3.54 ^c	0.084	-		
AL/GL ratio	0	1.20	0.99	1.04	1.67	1.88	0.073	1.35±0.032		
AL/GL Tatio	4	1.11	1.21	1.20	1.65		0.073	1.30±0.032		
Overall mea	n	1.15 [⊳]	1.10 ^b	1.12 [⊳]	1.66 ^a	1.61 ^a	0.051	-		
Urea	0	30.17	32.06			23.71		26.57 [⊳] ±0.555		
(mg/dl)	4	39.16	37.34		28.46	27.83	1.24	32.45 [°] ±0.555		
Overall mea	n	34.67 ^a	34.70 ^a	26.84 ^b	25.59 [⊳]	25.77 ^b	0.878	-		
AST	0	22.59	23.49	22.77	20.76	22.19		22.36 ^b ±0.232		
(U/L)	4	25.52	26.11	23.83	23.46	23.81	0.519	24.55 ^a ±0.232		
Overall mea	n	24.06 ^{ab}	24.80 ^a	23.30 ^b	22.11 ^c	23.00 ^b	0.367	-		
ALT	0	4.47	4.55	4.52	4.33	4.71	0.330	4.51 ^b ±0.147		
(U/L)	4	5.95	6.31	5.98	4.91	5.91	0.330	5.81 ^a ±0.147		
Overall mea	n	5.21 ^{ab}	5.43 ^a	5.25 ^{ab}	4.62 ^b	5.31 ^{ab}	0.233	-		

Table (10): Effect of experimental diet on biochemicals and enzyme activity in blood serum of goats.

Means with different litters within each row or column are significantly different (P≤0.05).

Milk yield and composition:

Data in Table (11) showed that urea treatments significantly (P \leq 0.05) increased milk yield, 4% fat corrected milk and milk composition more than control and untreated group. T4 had the highest (P \leq 0.05) values of milk yield, 4% fat corrected milk, and percentages of total solid, fat, solids not fat, total proteins, lactose and ash, followed by T5 then T3, while T2 and T1 had the lowest (P \leq 0.05) values. The difference among T4, T5 and T3 was significant

(P \leq 0.05), while the difference in ash content was not significant. Also, the differences between T2 and T1 were not significant for all values.

It is clear that, milk yield, 4% fat corrected milk and milk composition significantly (P \leq 0.05) increased by progressed stage of lactation. Only fat percentage increased (P \leq 0.05) at mid lactation stage then it decreased (P \leq 0.05) to reach the lowest value at late lactation stage. Total proteins concentration showed insignificant differences by progressed stage of lactation, but the differences between mid and late lactation was not significant.

These results have clearly demonstrated that sugar beet feeding to lactating goats did not have any adverse effects on milk production or milk composition. The increase in milk production by urea treatment may be attributed to the increase in feed intake in ureated treatments or may be due to the improvement in nutrient digestibility especially CP, CF and it components and the improvement in rumen fermentation especially VFAs concentration. Also, the increase in milk yield was attributed to the increase in molar proportion of acetic in the rumen, and to the increase in serum total proteins and albumin concentration, which reflect the better nutritional management. The decrease of milk fat content by progressed stage of lactation may be attributed to the increase of milk yield because there is a negative relationship between milk fat percentage and milk yield. The increase in milk production might be due higher water and sugar content of sugar beet (Niazi et al., 2000). The increase in milk lactose might be due to the high concentration of NDF, especially pectin in sugar beet. Further, pectin fermentation produces less lactate and a higher ratio of acetate to propionate without affecting cellulose and emicelluloses digestion (Abedo 2006).

Similar results were obtained by Wanapat *et al.* (2000), who found that the combination of ureated rice straw and whole sugar cane at the ratio of 50:50 enhanced both DM intake (P<0.05), milk yield, and milk fat and protein percentages. Man and Wiktorsson (2001) reported that milk protein was not different but milk fat increased in lactating cows fed fresh rice straw treated by urea. Sanh *et al.* (2002) indicated that urea treatment for rice straw increased milk yield, however, milk composition did not change. Al-Busadah (2008) indicated that milk yield, percent of water, fat, total solids, solid not fat, casein, pH and ash significantly (P< 0.05) differed in milk collected from goats fed urea-treated rice straw as compared to control.

Wanapat *et al.* (2013) reported that urea-calcium hydroxide treated rice straw significantly (P<0.05) increased milk yield and milk composition of lactating dairy cows. Also, Sorathiya *et al.* (2015) found that milk yield, milk fat percentage and 4% FCM were increased slightly in buffaloes fed sugar beets; however, they were similar to control, in percentages of SNF, proteins and lactose.

composition during different lactation periods:									
ltem	Stage	Experimental diet				±SEM	Overall		
item		T1	T2	T3	T4	T5		mean	
Milk	Early	898.1	896.3	940.0	982.5	968.1		937.00 ^c ±5.239	
yield	Mid	907.5	903.1	941.9	1015.6	999.4		953.50 ^b ±5.239	
(ml/h/d)	Late	1002.5			1234.4			1107.3 ^a ±5.239	
Overall mean		936.0 ^ª	933.1 [°]	997.9 ^c	1077.5 ^ª	1051.7 ^⁰	6.763	-	
4% FCM (ml/h/d)	Early	838.5	833.2	890.1	947.7	918.7	11.02	885.7 ^c ±4.932	
	Mid	848.6	843.3	893.4	983.3	955.1	11.02	904.7 ^b ±4.932	
	Late	923.6	918.8		1174.5	1113.3	11.02	1032.6 ^ª ±4.932	
Overall mean		870.2 ^d	865.1 ^d	938.8 ^c	938.8°1035.21995.7 ^b 6.367		-		
Milk composition (%):									
Total solids	Early	12.34	12.31	12.48	12.99	12.48	0.026	12.52 ^c ±0.011	
	Mid	12.42	12.39	12.53	12.98	12.76	0.026		
	Late	12.47	12.44	12.56	13.06	12.71	0.026	12.65 ^a ±0.011	
Overall mean		12.41 ^d	12.38 ^d	12.53 [°]	13.01 ^ª	12.65 ^b	0.015		
Fat	Early	3.55	3.53	3.64	3.76	3.66	0.013	3.63 [°] ±0.006	
	Mid	3.56	3.55	3.65	3.78	3.70	0.013	3.65 ^a ±0.006	
	Late	3.47	3.45	3.52	3.67	3.58	0.013	3.54 ^c ±0.006	
Overall mean		3.53 [₫]	3.51 ^d	3.61 ^c	3.74 ^a	3.64 ^b	0.008	-	
Solids not fat	Early	8.78	8.78	8.84	9.23	8.82	0.028	8.89 ^c ±0.012	
	Mid	8.85	8.83	8.88	9.19	9.06	0.028	8.96 ^b ±0.012	
	Late	9.00	8.98	9.03	9.38	9.13	0.028	9.10 ^a ±0.012	
Overall mean		8.88 ^d	8.86 ^d	8.92 ^c	9.27 ^a	9.00 ^b	0.016	-	
Total protein	Early	3.77	3.78	3.90	4.02	3.94	0.008	3.88±0.003	
	Mid	3.77	3.77	3.88	3.99	3.91	0.008	3.86±0.003	
	Late	3.77	3.75	3.89	4.018	3.94	0.008	3.87±0.003	
Overall mean		3.77 ^d	3.77 ^d	3.89 ^c	4.01 ^a	3.93 ^b	0.004	-	
Lactose	Early	3.946	3.93	4.26	4.39	4.30	0.022	4.16 ^c ±0.010	
	Mid	4.23	4.22	4.43	4.53	4.50	0.022	4.38 ^b ±0.010	
	Late	4.41	4.40	4.63	4.82	4.71	0.022	4.59 ^a ±0.010	
Overall mean		4.19 ^d	4.18 ^d	4.44 ^c	4.58 ^a	4.50 ^b	0.012	-	
	Early	0.984	0.984	1.037	1.045	1.043	0.004	1.019 ^ª ±0.002	
Ash	Mid	0.982	0.981	1.007	1.015	1.008	0.004	0.998 ^b ±0.002	
	Late	0.976	0.970	1.007	1.014	1.009	0.004	0.995 ^b ±0.002	
Overall mean		0.981 ^⁵	0.978 ^b	1.01 ^a	1.02 ^a	1.02 ^a	0.002	-	

 Table (11): Effect of experimental treatments on milk yield and composition during different lactation periods:

Means with different litters within each row or column are significantly different (P≤0.05).

Kid performance:

Data indicated that urea treatments of goat does significantly increased ($P \le 0.05$) its birth weight more than control and untreated group. The highest ($P \le 0.05$) birth weight was for T4, followed by T5 then T3 with insignificant differences between T4 and T5, while the lowest ($P \le 0.05$) birth weight was for untreated group, followed by control with insignificant differences. It seems that the body weight of kids during the first, second and third months (weaning weight) showed the same trend, but urea treatments had the highest ($P \le 0.05$) birth weights.

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2nd month

3rd month

verall mean

118.22

104.99^{ab}

105.55^c

121.40

102.77^b

105.80^c

Data of growth rate and overall mean of growth rate indicated that T4 had the highest values, followed by T5 then T3 with insignificant differences, except for the first month, while the lowest (P≤0.05) values were for untreated group, followed by control with insignificant differences in all values. These results of body weight and growth rate reflect the better nutritional management for goat dams fed SBP ureated with 2, 4 and 6%. The increase of body weight and growth rate in kids of goats fed ureated SBP can be fairly attributed to the higher feed intake than those fed CFM.

w	eight and	growth r	ate of thei	r kids.					
Item	Experimental diet								
item	T1	T2	Т3	T4	T5	±SEM			
Body weight									
Birth weight	2.47 ^{bc}	2.36 ^c	2.50 ^b	2.66 ^a	2.62 ^a	0.041			
1 st month	5.27 [°]	5.16 ^c	5.48 ^D	5.91 ^a	5.72 ^a	0.067			
2 ^{na} month	8.82 ^c	8.80 ^c	9.09 ^b	9.39 ^a	9.32 ^a	0.057			
/eaning weight	11.97 ^c	11.88 ^c	12.39 [⊳]	12.69 ^a	12.58 ^{ab}	0.077			
Growth rate (g/h/d):									
1 st month	93.44 ^c	93.22 ^c	93.88 ^c	113.52 ^a	103.33 ^b	2.931			

Table (12): Effect of feeding goat does the experimental diets on body

108.06^{bc} Means with different litters within each row or column are significantly different (P≤0.05).

120.29

110.00^a

116.10

110.00^a

113.20^a

119.92

108.51^a

110.59^{ab}

119.92

1.839

0.990

These results were supported by those obtained by El-Badawi and El-Kady (2006), who noted that inclusion of SBP spraved with urea solution at a level of 30 g/kg and replaced at 50% of the common CFM increased average daily gain by about 30%. Also, Gado et al. (2007) and Aziz (2009) reported an increase in body weight, average daily gain (ADG) and growth rate in sheep fed diets treated with 4% urea. Moreover, Okab et al. (2012) indicated that diet containing SBP treated with 4% urea improved (P<0.05) final body weight (FBW) and ADG of lambs. However, FBW and ADG decreased (P<0.05) in lambs fed on untreated diet compared to control diet. On the other hand, Bodas et al. (2007) and Omer et al. (2013) reported that inclusion of untreated SBP in sheep diet had no significant effect (P<0.05) on FBW, total body weight gain, and ADG.

CONCLUSION

It is clear that treating SBP at a level of 4% urea (T4) enhanced digestibility coefficients of all nutrients, fiber fractions, nitrogen balance, rumen parameters concentrations especially microbial protein concentration, ruminal protozoa count, serum total proteins and albumin, and milk yield and composition more than those treated at levels of 2 or 6% urea (T3 and T5, respectively). Therefore, the present study suggests the replacement of dried sugar beet pulp treated with 4% urea by a part of common concentrate feed mixture of lactating goats.

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تأثير تفل بنجر السكر المعامل باليوريا على أداء الماعز الحلابة هند أحمد عزيز 'و عبد القادر محمود خليف ' ١- قسم تغذية الحيوان والدواجن- مركز بحوث الصحراء-القاهرة- مصر. ٢- قسم الألبان - شعبة الصناعات الغذائية - المركز القومي للبحوث.

تمت هذه الدراسة على عدد ٤٥ من أناث الماعز البرقي لدراسة تأثير تفل بنجر السكر المعامل ب ٢، ٤و ٦% يوريا ليحل محل ٣٠ % من مخلوط المركزات الشائع في علائق الماعز على أداء الماعز الحلابة - محصول - تركيب اللبن - وزن الميلاد والفطام. تم تصميم خمس علائق يجريبيه كالتالي: معاملة (١): مخلوط مركزات + دريس برسيم (مقارنة). معاملة (٢): مخلوط مركزات+ تفل بنجر السكر غير معامل + دريس برسيم. معاملة (٣): مخلوط مركزات+ تفل بنجر السكر معامل ب ٢% يوريا (على أساس المادة الجافة) + دريس برسيم. معاملة (٤): مخلوط مركزات+ تفل بنجر السكر معامل ب ٤% يوريا (على أساس المادة الجافة) + دريس برسيم. معاملة (٥): مخلوط مركزات+ تفل بنجر السكر معامل ب ٦% يوريا (على أساس المادة الجافة) + دريس برسيم. أستمرت التجربة لمدة 140 يوم. وفي نهاية مرحلة الحليب تم اجراء تجربة هضم لدراسة تأثير التغذية بالعلائق التجربية على معاملات الهضم ، تخمرات الكرش، البروتين الميكروبي، بروتوزوا الكرش ومكونات الدم. أظهرت النتائج المتحصل عليها أن المعاملات الخامسة والرابعة والثالثة أدت إلى زيادة محتوى كلا من المادة الجافة، المادة العضوية، الدهن، المستخلص الخالي من الدهن والبروتين الخام، بينما أدت إلى نقص محتوى كل من الألياف الخام ومكوناتها، السليلوز والهيميسليلوز مقارنة بالمجموعة الأولى والثانية. وأظهرت المجموعة الرابعة والخامسة أعلى معاملات هضم لكل من المادة الجافة، المادة العضوية، الدهن، المستخلص الخالي من الدهن، البروتين الخام، الألياف الخام ومكوناتها، السليلوز والهيميسليلوز، كما أظهرت المجموعة الرابعة والخامسة أيضاً أعلى قيمة للمواد الكلية المهضومة والبروتين الخام المهضوم. أدت المعاملات الرابعة والخامسة والثالثة، على التوالي إلى تحسن معنوى في ميزان النيتروجين والماء وتخمرات الكرش وأعداد بروتوزوا الكرش ومحصول ومكونات اللبن ووزن الميلاد والفطام. و يتضح مما سبق أن معاملة تفل بنجر السكر باليوريا (٣%) له تأثير ملحوظ على أداء الماعز الحلابة و صغار ها.