

## **THERMOREGULATORY RESPONSES OF SHEEP TO STARVATION AND HEAT STRESS CONDITIONS**

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### **SUMMARY**

*The present study was carried out at Sakha Animal Production Experimental Station belonging to Animal Production Research Institute. The experiment was conducted under thermoneutral conditions (TNZ) in winter (THI < 74) and during moderate heat stress (MHS, THI 74 - <88) in summer season on 9 and 6 Ossimi rams, respectively. Body weight, thermoregulatory parameters and blood parameters were determined at the start of the experiment and after 48 and 96 hrs of starvation in each experiment. Ambient temperature (Ta), relative humidity (RH %) and Temperature-Humidity Index (THI) were measured simultaneously during the experiment.*

*Under TNZ, starvation had no significant effect on rectal temperature (RT), due to a significant and insignificant reduction in heat production (HP) and evaporative cooling (minute ventilation, MV), respectively. While under MHS, starvation reduced the effect of heat stress on RT by decreasing MV and respiration rate (RR), while, HP remained almost unchanged. Under TNZ, starvation caused an insignificant and significant decrease in plasma total cholesterol (Tcho.) and total lipids (TL), respectively. Meanwhile under MHS, starvation had no significant effect on plasma Tcho. and TL concentrations.*

*Before starvation, moderate heat stress caused a significant increase in RT; meanwhile, respiratory evaporation (MV) and heat production (HP) were lower than that in the TNZ. Moderate heat stress (MHS) did not affect significantly blood hematocrit (Ht) and plasma proteins (plasma total proteins (Tp), albumin (Alb) and globulin (Gl)) while it increased significantly plasma glucose concentration.*

*It could be concluded that starved sheep can tolerate moderate heat stress by decreasing evaporative cooling; meanwhile normal fed sheep depends on heat storage to tolerate moderate heat stress.*

**Keywords:** *Sheep, starvation, heat stress, thermoregulation, metabolic rate*

### **INTRODUCTION**

Ossimi sheep is an Egyptian native fat-tailed breed. Salem *et al.* (1982) reported that the Egyptian native fat-tailed sheep were shown to be more heat tolerant than the European breeds as fat-tailed sheep have lower rectal temperature and respiration rate than the European breeds under the same management and environmental conditions. Devendra (1982) stated that 43.3% of the total world populations of sheep are presented in tropical and subtropical regions. Sheep plays underestimated but

valuable role in the supply of dietary animal proteins and also socioeconomic status of several million small farmers, peasants and landless agricultural laborers.

Under Egyptian conditions, sheep, goats and camels are the main species for animal production in the desert areas. The usual watering and feeding in the desert conditions varies from one day to many days depending on the season, availability of water, feed and distance traveled between watering points. Under these conditions animals are faced with two physiological problems; namely, obtaining sufficient feed and water, and regulating their body temperature. The lack of feed and water would impose starvation and dehydration which affects productivity (body weight, fertility and milk production), thermoregulation (physiological reactions) and blood chemistry and metabolites (More *et al.*, 1983).

In tropical and subtropical regions animal are subjected to heat stress due to high environmental temperature and relative humidity during hot season. Heat stress affects significantly the heat balance of homeothermic animals and the main thermoregulatory mechanisms are the reduction in heat production and increase in heat loss (Johnson *et al.*, 2003). Khalifa *et al.* (1987), Khalifa *et al.* (2002), Srikandakumar *et al.* (2003) and Al-Haidary (2004) found that heat stress caused a significantly increase in rectal temperature and respiration rate of sheep and the effect was more pronounced in less adapted breeds (Srikandakumar *et al.* 2003). Mitlöhnnera *et al.* (2001) stated that during heat stress the coping strategy of cattle is to decrease metabolic heat production by lowering feed intake, which adversely affects productivity. Many authors reported that heat production decreased under heat stress (Luiting *et al.*, 1985 and Berman, 2003).

Bell *et al.* (1983) revealed that during heat stress, panting animals increase respiratory heat loss by increasing minute ventilation and when the heat stress is severe, the increase in minute ventilation includes an increase in alveolar ventilation and consequently, arterial CO<sub>2</sub> tension (PaCO<sub>2</sub>) declines resulting in a respiratory alkalosis.

Regarding the effect of starvation on thermoregulation, Brockway *et al.* (1965) found that starvation for 3 to 4 days decreased total heat loss, sensible heat loss and respiratory and skin evaporation of Cheviot sheep. Also, it has been found that fasting or starvation decreased VO<sub>2</sub>, VCO<sub>2</sub>, RQ and heat production (Blaxter and Waiman, 1966, Bennet, 1972, Kelly *et al.*, 1993 and Khalifa *et al.*, 2003).

The objectives of the present study were to determine the influence of feed deprivation under thermoneutral and moderate heat stress conditions on thermoregulation of native Egyptian sheep adapted to heat stress as well as to investigate the thermoregulatory mechanisms of normal and starved sheep to tolerate moderate heat stress under Egyptian conditions.

## MATERIALS AND METHODS

The present study was carried out at Sakha Animal Production Experimental Station belonging to Animal Production Research Institute. Fifteen unshorn (wool length more than 5 cm) Ossimi Rams (9 in winter season and 6 in summer season) were selected randomly and used in these studies. They were 2.5 to 3.0 years of age. All animals were fed concentrate and hay ration according to their requirements (NRC, 1985). The feed concentrate mixture contained 12% crude protein and 50% starch value. It was consisted of 35% corticated cotton seed meal, 22% corn, 33%

wheat bran, 4% rice bran, 3% molasses, 2% lime stone and 1% salt. Fresh tap water was available *ad lib*. Animals were kept indoors in metabolic cages during the experimental period to determine nitrogen balance. Thermoregulatory and blood parameters were determined immediately before (0 time) and after 48 and 96 hrs of starvation.

Ambient temperature (Ta) and relative humidity (RH %) were recorded simultaneously while measuring the physiological responses. The Temperature-Humidity Index (THI) was calculated from Ta and RH according to Thom (1959) converted to °C as follows:

$$\text{THI} = 9/5 * ((T*17.778)-(0.55-(0.55*RH/100))*(T-14.444)).$$

Where:

T = Dry bulb temperature in °C.

RH= Relative humidity as %.

The experiment was carried out under thermoneutral conditions (THI < 72) in winter (according to Fuquay (1981) and Khalifa (2005) and during mild to moderate heat stress (THI 76 – 78.5) in summer season (moderate heat stress according to Fuquay (1981) and mild to moderate heat stress according to Khalifa (2005).

In both seasons, the physiological parameters were measured every 48 hrs. at midday of the highest Ta from 12.00 to 14.00. Rectal temperature (RT, °C) was determined using a clinical thermometer (0.1 °C accuracy inserted 5 cm in the rectal for 1 min.). Skin temperature (ST, °C) and ear temperature (ET, °C) were measured using the Minolta/Land Cyclops Compac 3, a portable infrared thermometer (0.95% error and 0.1 °C accuracy). Respiration rate (RR) was determined by counting the flank movements in one minute. Respiratory minute volume (MV as l/minute) was measured by Dry Gas Meter. Tidal volume (TV) was calculated by dividing MV/RR. Heat production (HP) (measured as fasting metabolic rate, kcal/BW<sup>0.75</sup> per day) was calculated using the equation of Brouwer (1965). The measurement of oxygen consumption (VO<sub>2</sub>) and carbon dioxide production (VCO<sub>2</sub>) were made using the open-circuit technique according to Yousef and Dill (1969). Oxygen consumption was calculated from the oxygen deficit in expired air using oxygen analyzer (Servomex 570). The rate of carbon dioxide production was calculated from the CO<sub>2</sub> deficit in expired air obtained from infrared Gas Analyzer (Model-AR-411). Calculation of % true VO<sub>2</sub>, VCO<sub>2</sub> and volume VO<sub>2</sub> consumption and O<sub>2</sub> and CO<sub>2</sub> production and Respiratory Quotient (RQ) were done using the equations of Consolazio *et al.* (1963) where RQ= volume of CO<sub>2</sub> produced/volume of O<sub>2</sub> consumed.

Blood samples were collected from the jugular vein in heparinized tubes. Hemoglobin concentration (Hb, g/l) was determined by colorimetric methods using hemoglobin kits (Drabkins reagent) and hematocrite value (Ht, %) using microhematocrite technique. Plasma total protein (TP, mg/dl) and plasma albumin (Alb, mg/dl) concentrations were determined using the kits of the Egyptian American Co. Plasma globulin (Gl, mg/dl) was calculated by subtraction. Glucose concentration (Glu, mg/dl), total cholesterol (Tcho, mg/dl), and total lipids (TL, g/dl) were measured by colorimetric method using Sentinal kits.

The statistical analysis of data was carried out using SAS program (SAS, 1990). Proc GLM of SAS (two way analysis of variance with one way repeated measurement) was used to test the effect of heat stress, starvation and their interaction. To test the effect of interaction, analysis of variance and least square

means (Proc. LS means of SAS) were used to test the effect of starvation within each temperature group and Student t test (Proc MEANS of SAS) was used to test the effect of temperature within each starvation interval.

## RESULTS AND DISCUSSION

Meteorological data (Table, 1) reveal that mean ambient temperature ( $T_a$ ) and relative humidity during the experimental period ranged between 23.7 to 26.2 °C and 36 to 40% in the first experiment (in winter) and 27.9 to 33.7 °C and 26 to 54 % in the second experiment (in summer), respectively. The mean THI ranged between 68.4 to 71.9 and 76 to 78.5 in the first and second experiments, respectively. According to Fuquay (1981) and Khalifa (2005) sheep were under the thermoneutral zone range in winter and mild to moderate heat stress in summer.

**Table 1. Ambient temperature (AT°C), relative humidity (RH %) and Temperature-Humidity Index (THI) during the experiment period**

Parameter	Thermoneutral zone (TNZ)			Moderate heat stress (MHS)		
	0 time	48 hrs	96 hrs	0 time	48 hrs	96 hrs
AT	24.1	26.7	23.4	33.7	27.9	30.0
RH	40.0	38.0	35.9	25.7	54.0	39.8
THI	69.7	71.9	68.4	78.5	76.0	76.7

Fuquay (1981) stated that a THI of 72 and below is considered as no heat stress, 73-77 as mild heat stress, 78-89 as moderate and above 90 as severe, while Khalifa (2005), based on changes in body temperature, heat production and heat loss of sheep, reported that the comfort zone ranged between 20-25 °C (60-70 THI) while moderate heat stress ranged from 25 – 28 °C (70 – 85 THI) and severe heat stress occurred at  $T_a \geq 28$  (THI  $\geq 85$ ).

### *Effect of ambient temperature*

At zero time (before starvation), moderate heat stress (MHS) caused a significant increase in RT; meanwhile, respiratory evaporation (MV) and heat production (HP) were lower than in the TNZ. The significantly lower respiratory evaporation under MHS than in TNZ was due to an insignificant decrease in RR and TV as an adaptive mechanism to compensate the reduction in HP under heat stress (Table 2). Similar trends were found after starvation for 48 hrs where RT was significantly higher under MHS than under TNZ, meanwhile, respiratory evaporation (RR, TV and MV) and gas exchange ( $VO_2$  and  $VCO_2$ ) were significantly and HP was insignificantly lower under MHS than under TNZ. However, after starvation for 96 hrs, MHS had no significant effect on RT and HP, while respiratory evaporative (RR and MV) was significantly lower under MHS than under TNZ.

Khalifa *et al.* (2002) and Al-Haidary (2004) reported that heat stress significantly increased RT, RR and ST of sheep, while, Ismail *et al.* (2002) found that RT did not differ significantly between summer and winter. Meanwhile, Butswat *et al.* (2000) stated that an increase in respiration rate appeared to be the immediate response of sheep to environmental stress. The effect of heat stress on RT and RR depends on the severity of heat stress as indicated by Lowe *et al.* (2002) who reported that RR and

RT were highly correlated with increasing THI. The insignificant changes in RR under MHS is in accordance with Srikandakumar *et al.* (2003) who concluded that the higher magnitude of increase in RT and lower magnitude of increase in RR in Omani sheep (local adapted breed) than in Merino ones (exogenous breed) during the period of heat stress suggests that this breed was less stressed than the Merino sheep with increasing heat stress. They suggested that this adapted breed of sheep can store body heat during the periods of heat stress, which can economize on the loss of water and the increased need of energy under these conditions.

Higher ST and lower ET under TNZ than under MHS (Table, 2) may correspond to vasodilatation under heat stress (Turnpenny *et al.*, 1997). Khalifa (1982) and Al-Haidary (2004) reported that heat stress significantly increased ST of sheep; however, Khalifa (1982) found that ET was also significantly higher in summer than in winter.

The reduction in TV under MHS is an adaptive mechanism to prevent respiratory alkalosis. Bell *et al.* (1983) revealed that during heat stress, panting animals increase respiratory heat loss by increasing minute ventilation (MV). When the heat stress is severe, the increase in MV includes an increase in alveolar ventilation, and consequently, arterial CO<sub>2</sub> tension declines resulting in a respiratory alkalosis.

The insignificant decrease in HP under MHS is an adaptive mechanism to minimize the disturbance in heat balance indicated by significant increase in RT. Many authors reported that HP decreased under heat stress (Luiting *et al.*, 1985 and Berman, 2003).

Moderate heat stress had no significant effect on RQ although the significant reduction in both of VO<sub>2</sub> and VCO<sub>2</sub> due to the reduction in TV under MHS (Table 2) which confirm the previous conclusion that the reduction in TV under MHS is an adaptive mechanism to prevent respiratory alkalosis.

Heat stress had no significant effect on Ht value. However, Hb was significantly lower under MHS than in the TNZ but the difference was insignificant after starvation for 96 hrs due to a significant reduction in Hb during starvation under TNZ (Table, 3). Srikandakumar *et al.* (2003) reported that heat stress decreased ( $P<0.01$ ) Hb in the Merino sheep. Meanwhile, Al-Haidary (2004) reported that heat stress significantly increased Ht of Naimey sheep. Khalifa *et al.* (1987) indicated that the increase in Ht and Hb of sheep after exposure to solar radiation was due to the reduction in plasma volume. Accordingly, the decrease in Hb in the present results reveals that MHS did not affect plasma volume. The insignificant effect of heat stress on Hb after starvation for 96 hrs. is in accordance with Cole and Hutcheson (1985) who indicated that the degree of changes in Ht and Hb were more pronounced in ad libitum animals than in the feed-restricted animals.

Plasma TP, Alb and Gl concentrations did not differ significantly between TNZ and MHS except after starvation for 96 hrs when plasma TP and Alb were significantly lower under MHS than under TNZ due to a significant increase in TP and Alb after starvation for 96 hrs under TNZ. Macfarlane *et al.* (1959 and 1966) reported that plasma total protein of sheep changed little between summer and winter. However, Ismail *et al.* (2002) found that TP and Gl were higher in summer than in winter.



**Table 3. Blood parameter of Ossimi sheep as affected by Thermoneutral zone (TNZ), Moderate heat stress (MHS) and starvation conditions**

Parameter	Thermoneutral zone (TNZ)				Moderate heat stress (MHS)				P		
	0 time	48 hr	96 hr	S.E.	0 time	48 hr	96 hr	S.E.	0 time	48 hr	96 hr
Ht %	31.4	29.9	29.8	1.09	30.2	32.1	34.0	1.64	NS	NS	NS
Hb (g/dl)	19.1 <sup>a</sup>	20.5 <sup>a</sup>	13.1 <sup>b</sup>	0.71	11.3	12.9	12.5	0.94	**	**	NS
Glu. (g/dl)	17.0	14.3	25.8	4.57	25.2 <sup>b</sup>	44.0 <sup>ab</sup>	63.8 <sup>a</sup>	12.3	**	**	NS
TP (g/dl)	6.71 <sup>b</sup>	7.87 <sup>b</sup>	10.47 <sup>a</sup>	0.53	7.78	7.52	7.66	0.39	NS	NS	*
Alb (g/dl)	3.06 <sup>b</sup>	4.03 <sup>a</sup>	4.11 <sup>a</sup>	0.26	3.44	2.54	2.68	0.34	NS	**	**
Gl (g/dl)	3.65 <sup>b</sup>	3.84 <sup>b</sup>	6.36 <sup>a</sup>	0.53	4.34	4.98	4.98	0.51	NS	NS	NS
Tcho. (mg/dl)	86.7	76.6	81.2	8.2	34.8	43.9	50.8	8.7	**	**	*
TL (g/dl)	4.643 <sup>a</sup>	3.253 <sup>b</sup>	3.377 <sup>b</sup>	0.32	2.68	2.51	2.82	0.29	**	**	**

a,b,c similar letters within each column and variable are not significant ( $p \leq 0.05$ ).

p = Probability level for the effect of ambient temperature.

SE = Standard error of the LEAST square means.

NS = Insignificant  $p > 0.05$ . \* = Significant at  $p \leq 0.05$ . \*\* = Significant at  $p \leq 0.01$ .

Hbg/l = Hemoglobin concentration.

Ht % = Hematocrite value.

TP mg/dl = Plasma total protein.

Alb, mg/dl = Plasma albumin.

Gl mg/dl = Plasma globulin

Glu mg/dl = Glucose concentration

Tcho mg/dl = Total cholesterol

TL g/dl = Total lipids.

Plasma Glu concentration was significantly higher under MHS than under TNZ either before or after starvation for 96 hrs. Similar results were found by Samak *et al.* (1986) but contradict with those of Ismail *et al.* (2002) who reported that Glu was significantly lower in summer than in winter. Seasonal changes in Glu depend on changes in RR and HP (Shaffer *et al.*, 1981). Opposite trends were found in plasma Tcho. and TL where they were significantly lower under MHS than under TNZ either before or after starvation (Table 3). Similar results were found by Ismail *et al.* (2002). The significant increase in plasma Glu concentration and decrease in plasma Tcho. and TL under MHS agree with Srikandakumar *et al.* (2003) who found that heat stress increased ( $P < 0.01$ ) plasma glucose in Merino (less adapted) but decreased ( $P < 0.01$ ) it in Omani (more adapted) sheep. Nazifi and Gheisari (1999) found that the concentrations of serum cholesterol, triglyceride, total lipid, HDL-cholesterol, LDL-cholesterol and VLDL-cholesterol of dromedary camels were significantly higher ( $p < 0.05$ ) in winter months than in summer months. Results showed that very hot and cold conditions had a considerable effect on serum lipids in dromedaries.

The present results indicate that sheep can tolerate MHS by decreasing HP and increasing heat storage (significant increase in RT) to compensate the significant reduction in evaporative cooling (due to a decrease in TV and RR) as an adaptive mechanisms to conserve water under natural MHS. Folk (1974) stated that body temperature is naturally maintained at a relatively constant level because of the balance which exists between heat production and heat loss. Starved sheep can tolerate moderate heat stress by decreasing evaporative cooling, while normal fed sheep depend on heat storage and decrease of HP to tolerate MHS.

#### **Effect of starvation**

Starvation for 96 hrs caused a significant reduction in body weight by about 4% from the initial body weight in both seasons.

Under TNZ, starvation had no significant effect on RT, but ST and ET decreased significantly which was accompanied with significant reduction in MV due mainly to

significant reduction in RR. The insignificant effect of starvation on RT was due to the significant reduction in both of evaporative cooling (MV) and HP (Table 2).

Under MHS, RT decreased significantly after starvation for 4 days which was accompanied with a significant reduction in evaporation due to significant reduction in RR, meanwhile starvation had no significant effect on TV and HP. On the other hand under TNZ, starvation affected significantly thermoregulation by decreasing HP and evaporative heat loss (Table 2).

The significant reduction in RT and RR after starvation for 4 days without significant effect on TV and HP indicate that starvation under MHS prevented the effect of HS on RT so that MV and RR decreased, while, HP remained almost unchanged. Piccione *et al.* (2002) stated that prolonged food deprivation is known to cause a fall in the core body temperature of homeotherms. Li *et al.* (2000) found that fasting decreased HP and RT ( $P < 0.05$ ) and Ahmed and Abdelatif (1994) stated that RT and RR were reduced by food restriction. Meanwhile, Khalifa *et al.* (1999a and b) found that starvation increased RT and decreased RR in both shorn and unshorn ewes under cold conditions. They explained that the significant reduction in RR of ewes in response to 2 days of starvation may be a thermoregulatory mechanism to decrease heat dissipation through panting. Finch and King (1982) attributed the reduction in RT during restriction of feed (50% of maintenance) for 3 months to the reduction in metabolic rate and that the lower metabolic demands indicate that energy reserves could be spared from rapid depletion as starvation advances.

Under TNZ, starvation caused a significant reduction in  $VO_2$ ,  $VCO_2$  and HP (Table 2). While under MHS, starvation had no significant effect, but RQ and HP tended to decrease after starvation. Under both temperatures RQ tended to decrease with starvation due to more resuction in  $VCO_2$  than in  $VO_2$  under TNZ and an insignificant reduction in  $VCO_2$  with slight increase in  $VO_2$  under MHS. The reduction in RQ during starvation under TNZ and MHS, although it was insignificant, is in accordance with Bennet (1972) and Brockway *et al.* (1965) who found that the RQ declined significantly in sheep as the duration of fasting increased. Kelly *et al.* (1993) added that  $O_2$  consumption of ewes fed twice maintenance was higher than that of those fed maintenance ration or starved. Khalifa *et al.* (1999a and b) revealed that heat production, as indicated by  $VO_2$  and  $VCO_2$  changes, was significantly ( $P < 0.05$ ) decreased by starvation and duration of starvation in both shorn and unshorn ewes, reaching the lowest values at 4 days of starvation. This decrease in HP was due to the reduction in both metabolic heat production and specific dynamic action of food due to starvation. They also demonstrated that the reduction in RQ indicates that starvation caused more reduction in  $VCO_2$  than in  $VO_2$  due to catabolism of protein (RQ near 0.8) and fat (0.7) instead of mixed diet or carbohydrate (almost 1). As a result, after 2 days of food deprivation both shorn and unshorn ewes began to use the storage fat and body proteins as sources of energy.

Under both climatic conditions (TNZ and MHS) starvation had no significant effect on Ht value. Similar result was found in Hb under moderate heat stress, while under TNZ a significant reduction in Hb occurred after 96 hrs of starvation (Table 3). Ali *et al.* (1984) reported that feed restriction for 96 hr or 168 hr in male Nubian goats caused a decrease in Hb concentration, Ht and erythrocytes number. Panaretto (1964) added that undernourished decreased haematocrit value by decreasing red cell volume while plasma volume remained unchanged. Under cold conditions, Khalifa *et al.* (1999 a, b) detected a significant ( $P < 0.05$ ) decrease in the Hb concentration

accompanied with a non significant decrease in the Ht value in the shorn and unshorn ewes at day 4 of starvation, which indicating anemia.

Starvation had no significant effect on plasma proteins (TP, Alb and Gl) except a significant increase in TP after starvation for 96 hrs under TNZ due to significant increase in Alb and Gl indicating that under TNZ starvation may cause a reduction in plasma volume. Similar results were found by Vihan and Rai (1984) who reported that TP values were significantly reduced in 5 days starved sheep and goats.

Under MHS starvation increased significantly plasma Glu as a source of energy to maintain HP at the pre-starvation level. While under TNZ, an insignificant increase in plasma glucose occurred after 96 hrs of starvation (Table, 3). Rule *et al.* (1985) found that plasma glucose concentration in steers increased from day 2 to day 5 of fasting where it remained at this level up to day 8 of fasting. The increase in plasma glucose level during starvation may be due to the increase in growth hormone during starvation which causes the increase in glucose by its effect on gluconeogenesis. Martin (1976) stated that growth hormone secretion increases when plasma glucose levels fall during limited periods of food deprivation. The hormone promotes glucose formation from hepatic glycogen and pyruvate; it also modifies glucogenic influences of glucocorticoids.

Under TNZ, starvation caused an insignificant and significant decrease in plasma Tcho. and TI, respectively. Meanwhile, under MHS starvation had no significant effect on plasma Tcho. and TI concentrations although plasma Tcho. Tended to decrease insignificantly during starvation (Table 3). The reduction in plasma Tcho. and TI during starvation under TNZ disagrees with previous results of Khalifa *et al* (1986) who found that starvation increased significantly plasma free fatty acids and triglycerides of both shorn and unshorn ewes, while it had no significant effect on TL. Also, Cole and Hutcheson (1985 &1988) found that serum cholesterol was not significantly affected by feed deprivation and that serum cholesterol levels were negatively related to the feed intake of the steers.

## CONCLUSIONS

The main thermoregulatory mechanisms of starved sheep under MHS are the reduction in evaporative cooling, while in normal fed sheep heat storage is the main thermoregulatory mechanism to tolerate MHS.

Sheep can tolerate MHS by decreasing HP and increasing heat storage (significant increase in RT) to compensate the significant reduction in evaporative cooling (due to a decrease in TV and RR) as an adaptive mechanisms to conserve water under natural moderate heat stress.

Under TNZ starvation had no significant effect on RT, due to a significant and insignificant reduction in heat production and evaporative cooling (MV), respectively. While under MHS, starvation reduced the effect of heat stress on RT by decreasing MV and RR, while, HP remained almost unchanged.

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## التنظيم الحرارى فى الاغنام المعرضة للتصويم والاجهاد الحرارى

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أجريت هذه الدراسة فى محطة بحوث الانتاج الحيوانى بسخا – محافظة كفر الشيخ- معهد بحوث الإنتاج الحيوانى بهدف دراسة تأثير كل من التصويم و الإجهاد الحرارى على التنظيم الحرارى فى كباش الأغنام الاوسيمى المصرية. تحت انسب مدى حرارى TNZ و الاجهاد الحرارى المتوسط MHS تمت الدراسة على عدد 15 كبش اوسيمى ( عدد 9 كباش فى فصل الشتاء TNZ حيث تراوحت درجة الحرارة بين 23.4 – 26.7° م و عدد 6 كباش فى فصل الصيف MHS حيث تراوحت درجة الحرارة بين 27.9- 33.7° م ). تم قياس كل من درجة حرارة الجو Ta و الرطوبة النسبية RH% و دليل الحرارة و الرطوبة THI كذلك تم تقدير وزن الجسم و مقاييس التنظيم الحرارى (درجة حرارة المستقيم RT و درجة حرارة الجلد ST و درجة حرارة الاذن ET و معدل التنفس RR و حجم التنفس فى الدقيقة MV و كمية الاوكسجين المستهلكة VO<sub>2</sub> و كمية ثانى اكسيد الكربون المنتجة VCO<sub>2</sub> و النسبة التنفسية RQ و الانتاج الحرارى HP). و كذلك تم أخذ عينات دم من الكباش لتقدير (الهيموجلوبين و الهيماتوكريت و البروتينات الكلية و الالبومين و الجلوبيولين و الجلوكوز و الكوليسترول و البيبيدات الكلية) وذلك فى بداية فترة النجوع و بعد 48 ساعة ثم بعد 96 ساعة و أوضحت أهم النتائج الآتى:

- 1- الميكانيكية الرئيسية للتنظيم الحرارى للأغنام المعرضة للتصويم تحت الإجهاد الحرارى هى خفض التبريد عن طريق التبخير فى حين أن هذه الميكانيكية فى الأغنام المغذاه هى تخزين الحرارة.
- 2- الأغنام المغذاه تحت الإجهاد الحرارى تقوم بخفض الإنتاج الحرارى و زيادة تخزين الحرارة (زيادة درجة حرارة الجسم) لمواجهة النقص المعنوي فى التبريد عن طريق التبخير من سطح الجهاز التنفسي نظرا لنقص كل من معدل النفس و عمق التنفس و ذلك للمحافظة على ميزان ماء الجسم.
- 3- التصويم يخفض من تأثير الإجهاد الحرارى على الأغنام عن طريق خفض معدل التنفس و كذلك حجم الأوكسجين المستهلك فى النفس دون تغيير الإنتاج الحرارى.



**Table 2. Thermoregulatory responses of Ossimi sheep under Thermoneutral zone (TNZ), Moderate heat stress (MHS) and starvation conditions**

parameter	Thermoneutral zone (TNZ)				Moderate heat stress (MHS)				P		
	0 time	48 hrs	96 hrs	S.E.	0 time	48 hrs	96 hrs	S.E.	0 time	48 hrs	96 hrs
RT °C	39.1	39.3	39.3	0.09	40.1 <sup>a</sup>	40.0 <sup>a</sup>	39.1 <sup>b</sup>	0.14	**	**	NS
ST °C	38.0 <sup>a</sup>	37.7 <sup>a</sup>	36.3 <sup>b</sup>	0.36	37.6	36.7	37.4	0.32	NS	**	NS
ET °C	36.3 <sup>a</sup>	33.4 <sup>b</sup>	32.7 <sup>b</sup>	0.81	37.1 <sup>a</sup>	32.6 <sup>b</sup>	33.0 <sup>b</sup>	0.34	NS	NS	NS
RR (r/min.)	45.1	34.7	39.6	4.0	41.7 <sup>a</sup>	24.0 <sup>b</sup>	23.0 <sup>b</sup>	2.8	NS	*	**
TV (ml/breath)	120 <sup>ab</sup>	157 <sup>a</sup>	104 <sup>b</sup>	14.4	103	102	116	9.8	NS	*	NS
MV (l/min.)	4.529	4.575	3.461	0.39	3.278 <sup>a</sup>	2.121 <sup>b</sup>	2.254 <sup>b</sup>	0.20	**	**	**
VO <sub>2</sub> L/d.BW <sup>0.75</sup>	5.037 <sup>a</sup>	5.327 <sup>a</sup>	3.271 <sup>b</sup>	0.87	3.725	3.725	3.808	0.36	**	**	NS
VCO <sub>2</sub> L/d.BW <sup>0.75</sup>	4.629 <sup>a</sup>	4.477 <sup>a</sup>	2.841 <sup>b</sup>	0.71	3.791	2.966	3.152	0.48	**	**	NS
RQratio	0.92	0.84	0.89	0.06	1.02	0.80	0.83	0.08	NS	NS	NS
HP /day.BW <sup>0.75</sup>	24.2 <sup>a</sup>	25.4 <sup>a</sup>	16.6 <sup>b</sup>	2.9	18.0	18.0	18.9	1.74	NS	NS	NS

a,b,c similar letters within each column and variable are not significant (p≤0.05)

p = Probability level for the effect of ambient temperature

SE =Standard error of the LEAST square means.

NS = Insignificant p>0.05. \* = Significant at p≤0.05. \*\* = Significant at p≤0.01.

RT °C = Rectal temperature. ST°C = Skin temperature. ET °C = ear temperature.

RR r/min. = Respiration rate.

MV l/minute = Respiratory minute volume.

TV (ml/breath) = Tidal volume.

HP /day.BW<sup>0.75</sup> = Heat production.

VO<sub>2</sub>L/d.BW<sup>0.75</sup> = Oxygen consumption.

VCO<sub>2</sub>L/d.BW<sup>0.75</sup> = carbon dioxide production.

RQratio = Respiratory Quotient.