EFFECT OF THYROIDISM ON PRODUCTIVE PERFORMANCE OF GROWING NEW ZEALAND WHITE RABBITS UNDER HOT SUMMER CONDITION IN EGYPT

El-Sawy, M. A.1; Hoda A. Shabaan1 and A. A. Nada2

1- Animal Production Research Institute, Agricultural Research Center, Ministry of Agriculture, Dokki, Giza, Egypt.

2- Animal health Research Institute, Agricultural Research Center, Ministry of Agriculture, Dokki, Giza, Egypt.

ABSTRACT

Total of thirty growing New Zealand White rabbits (NZW) aged 35 days and 624±21.5(g) live body weight were used in this study. Rabbits were divided into similar three groups and treated once weekly as follows: The 1st group (control) was injected with saline solution, the 2nd and the 3rd groups were treated by eltroxin (hyperthyroidism) and carbimazole (hypothyroidism), respectively for two moths.

At 12 weeks of age, feed intake, body weight and body gain in carbimazole treatment were higher than eltroxin and control. Also, it improved growth rate and feed conversion than control by 8% and 3%, respectively. At 12 weeks of age, the minimum rectal temperature was found in eltroxin group.

Carbimazole had the higher final body weight than eltroxin and control by 11.1% and 13.5%, respectively. Dressing percentage was significantly (P<0.05) higher in eltroxin and carbimazole than that of the control group. However, weights of liver, kidneys, lungs and heart were almost higher in the control than both tested groups.

Level of T_4 and \tilde{T}_3 in control group increased compared to hyperthyroidism or hypothyroidism. There is no hematological change in both hyper and hypothyroid groups.

Keyword: Rabbit, thyroidism, body weight, T₃, T₄, carcass, rectal temperature.

INTRODUCTION

Rabbits exhibit seasonal patterns in production. In sheep, removal of the thyroid gland increased the production as reported by (Karsch et al., 1995). It is not known whether the thyroid gland plays a key role in the transition to production and reproduction in animals (DeMoraes et al., 1998). The thyroid gland has been known to regulate the metabolic activity in all animals (lossa et al., 2001). Thyroid hormones regulate basal metabolism and can alter nutrient requirements by increasing basal expenditure of energy. Induction of hypothyroidism in animals has been successfully accomplished using potent antithyroid compounds (Burroughs et al., 1960).

The hypothyroidism increases live body weight by improving body condition and feed efficiency when fed for short duration to steers (Oyedipe et al., 1982). Hypothyroidism decreased serum levels of thyroid hormones, which is due to decreased activity of the thyroid gland and the hypothalamus-pituitary

axis, (Thrift et al., 1999). Hyperthyroidism (thyrotoxicosis) is a multisystemic disorder resulting from excessive circulating concentrations of T_3 and/or T_4 (Thoday, 1993).

The objectives of this experiment were to evaluate the effects of hypo and hyper-thyroidism on productive performance of growing New Zealand White rabbits under summer season.

MATERIALS AND METHODS

This study was carried out during the period from August 2004 till September 2004 in a private rabbit's farm (Haraz farm) at Abou-El-Atta village, Nubaria, El-Behera Governorate. The analytic work was carried out in Animal Production and Animal health Research Institutes, Agricultural Research Center, Ministry of Agriculture, Dokki, Giza, Egypt.

Thirty growing New Zealand White rabbits (NZW) aging 35 days and weighing 624±21.5(g) live body weight was used in this study. Rabbits were randomly divided into three experimental groups (10 rabbits in each). Rabbits were reared in open rabbitry house system, and kept in individually wire cages for (35X45X30 cm). Rabbits were fed ad libitum on ration containing 17% crud protein and 2721 digestible energy (Kcal/kg) and clean fresh water was available by nipple drinker at all the time. All experimental animals were reared under the same summer conditions (26-31°C). Environmental temperature outdoors and indoors were recorded daily at (08.00, 14.00 and 22.00 h) to obtain the weeks average of ambient temperature as shown in Table (1).

Table (1): Means±SE of environmental air temperature outdoors and indoors.

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Temperature week	re Outdoors			Indoors		
	08.00h	14 <u>.</u> 00ħ	22.00h	08.00h	14.00h	22.00հ
1	30 86+0.40	35.00±0.62	25.14+0.51	26.14±0.67	30.57±0.43	29.14 <u>+</u> 0.51
3	31.00 <u>+</u> 0.31	36.12±0.53	23.71±0.42	26.57±0.57	31.17±0.37	29.43±0.43
5	30.14+0.80	34.00±0.53	27.00±0.44	27.43+0.57	28.71±0.29	28.71 <u>+</u> 0 57
7	26.43 <u>+1.19</u>	30.29+0.71	23.14+0.80	25.43±0.20	29.43±0.72	27.43 <u>+</u> 0.65

All experimental animals were injected subcutaneous 0.5 ml once weekly as the follows: the first group, was injected with saline solution (control), the second and third groups were injected 0.5 ml eltroxin (hyperthyroidism) and/or carbimazole (hypothyroidism) solution to provide 15 µg/rabbit of them. The rabbits were weighed at the beginning of the experiment and bi-weekly thereafter. Body weight and feed intake were recorded bi-weekly; then body gain and feed conversion were calculated and growth rate (%) was calculated by the following equation.

Blood samples were taken at 12 weeks old. Plasma samples were prepared by centrifugation (3000-rpm for 15 minutes) and stored at -20°C until analysis. Quantitative measurement of T_4 was carried by DSL-3200-USA ACTIVETM Thyroxin coated tube radioimmunoassay kits. The sensitivity, calculated by interpolation of mean minus two standard deviations of 12 replicates of the 0 µg/dl T_4 standard, is 0.4 µg/dl (Amer and Chicago, 1995). Quantitative measurement of T_3 was carried by DSL-3100-USA ACTIVETM Triiodothyronine coated tube radioimmunoassay kits. The sensitivity, calculated by interpolation of mean minus two standard deviations of 20 replicates of the 0 ng/dl T_3 standard, is 4.3 ng/dl (Tietiz, 1995). Carcass traits have been done for five rabbits from each group at 12 weeks old. Data were statistically analyzed using SAS (1995). Means were compared (P<0.05) using Duncan's multiple range test (Duncan, 1955) according the following model:

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

Where: Y_{ij} = observation of the ij^{th} rabbits, μ = overall mean, commn element to all observations, T_{i} = effect i^{th} group, e_{ij} = random error component assumed to be normally distribute.

RESULTS AND DISCUSSION

Growth performance:

Effects of treatments on daily feed intake (FI) and live body weight (LBW) in growing NZW rabbits are shown in Table (2). At 6 weeks old, rabbits treated with eltroxin or carbimazole, showed slight decrease in both feed intake and body weight as compared to the control, this observation is in agreement with the finding of Harada and Kato (1982). Concerning the group treated with eltroxin, the catabolic action of thyroxin could prevail without an increase in feed intake, leading to a decrease of the body weight (Dojana et al., 2000). This may be due to the reduction of thyroid hormone (hypothyroidism) caused to decrease of feed intake (Freake at al., 2001).

Although, rabbits treated with carbimazole and eltroxin slightly increased their feed intakes than the control group at 8 weeks old by 15.52% and 1.54%, respectively, they showed lighter body weight than the control group at the same old by 7.73% and 5.12%, respectively. However, hyperthyroidism showed decreasing in FI and increasing BW than hypothyroidism at 8 weeks old. This observation is in agreement with the finding of Marai et al. (1994). Thyroidectomized (hypothyroidism) may increased rate of body protein breakdown and decreased both plasma T_3 and protein synthesis or it is occur in proportion to changes in total body mass (Buttery and Lindsay, 1980).

Hypothyroidism caused increasing in feed intake and body weight than hyperthyroidism and control at 10 weeks old. This observation was agreement with the finding of LeGrow *et al.*, (1999). They suggesting that growth was minimal in the hyperthyroidism state. This result also, may be due to the increasing of catabolism of feedstuffs in hyperthyroidism (Thrift *et al.*, 1999). The differences in the results at all ages may due to that T_4 level diminished, reaching its nadir and it then increased to approximate its basal level (Shirpour *et al.*, 2003).

At 12 weeks of age, rabbits treated with carbimazole showed higher feed intake than eltroxin and control by 2.27% and 14.22% and higher in body weight by 9.9% and 4.9%, respectively. The increase in body weight with may be due to the fact that T_4 stimulates protein synthesis and increases each of nitrogen retention, sensitivity of tissues to GH (Ingbar and Woeber, 1981).

Table 2: Means+SE of daily feed intake (FI) and live body weight (LBW) of

growing NZW rabbits in different experimental groups.				
ltems	Control	Eltroxin	Carbimazole	
Feed intake (g):				
6 weeks	67.2 <u>+</u> 2.0	66.4 <u>+</u> 1.1	66.5 <u>+</u> 1.2	
8 weeks	75.1 <u>+</u> 2.2 ^b	76.3 <u>+</u> 1.2°	86.8 <u>+</u> 3.5°	
10 weeks	106.5±4.0 [▶]	114.5±2.7 ^{ab}	120.1±6.6 ^a	
12 weeks	143.2 <u>+</u> 7.7°	159.9 <u>+</u> 4.3 ^{ab}	163.5 <u>+</u> 17.2ª	
Live body weight (g):				
5 weeks (Initial)	641.5 <u>+</u> 9.1	583.3 <u>+</u> 27.7	622.2 <u>+</u> 27.8	
6 weeks	817.5 <u>+</u> 18.3	777.8 <u>+</u> 25.2	788.9 <u>+</u> 35.1	
8 weeks	1215±27.44°	1152.8 <u>+</u> 29.6 ^b	1121.1±44.9 ^b	
10 weeks	1585±39.5ab	1500±41.7 ^b	1652.8±52.1°	
12 weeks	1885±56.8b	1800 <u>+</u> 61.2 ^b	1977.8 <u>+67.2ª</u>	

Values having different superscripts in the same row are significantly different (P<0.05).

On the other hand, the hyperthyroidism decreased body weight at 8 and 12 weeks old by (-5.1% and - 4.5%) than control, respectively (Table 2). This case known by thyrotoxicosis which are a reflection of the stimulation of metabolism induced by an excess of thyroid hormones in circulation then weight loss (Baulieu and Kelly, 1990). This may be due to the concentration of receptors is affected by the level of hormone and this has been suggested as an especially important homeostatic regulatory mechanism in cell communication and function (Buttery and Lindsay, 1980).

Changes in rectal and skin temperatures (Table 3) differed among groups at different ages studied. The maximum rectal and skin temperatures were observed in eltroxin group at 8 weeks of age. This may be due to hyperthyroidism caused heat intolerance (Thoday, 1993). Also, the minimum rectal and skin temperatures were observed in carbimazole group at 8 weeks of

age. This observation is in agreement with the finding of Thrift et al. (1999). They reported that the decline in rectal temperature of hypothyroidism is indicative of a change in metabolism. This contributes to the increased basal metabolic rate (O2 consumption by the whole animal at rest) and increased sensitivity to heat in hyperthyroidism, and the converse in hypothyroidism (Greenspan and Bsxter, 1994). Rectal and skin temperatures of growing rabbits were differed significantly from normal values during the period of this experimental (Table 3). At 12 weeks of age, the maximum rectal temperature was found in carbimazole group. The increase in rectal temperature of hypothyroidism in the first days post-treatment may be correspond to the increase in thyroid hormone and the decline of it is indicative of a change in metabolism, DeMoraes et al. (1998). Hypothyroidism is characterized by a decreased rate of oxygen consumption per unit of body surface area and a decrease in heat production (Loeb, 1996a).

Table 3: Means±SE of rectal temperature (RT) and skin temperature (ST) in growing NZW rabbits in different experimental groups.

Treatment	Con	Control Eltro		xin Carbimazole		nazole
Age at	RT	ST	RT	st	RT	ST
5 weeks	39.97±0.09ª	39.2±0.08ª	39.69±0.11 ⁸⁶	39.09±0.12 ^b	39.59±0.13b	39.32±0.09ªb
6 weeks	40.00±0.17	39.00±0.15	40.78±0.08	39.18±0.08	39.86±0.08	38.81±0.08
8 weeks	39.95±0.13*	39.95±0.13	40.04±0.13 ^a	40.01±0.12	39.79±0.13 ^b	39.85±0.11
10 weeks	39.99±0.12 ⁸	39.98±0.13	39.94±0.10 ^a	39.86±0.08	39.82±0.05b	39.87±0.06
12 weeks	39 51±0.14°	39.46±0 13	39.26±0.09b	39.44±0.09	39.54±0.09a	39.40±0.08
* Values having different superscripts in the same row are significantly different (P<0.05).						

Body gain improved with eltroxin than control and carbimazole by 22.73 % and 16.63% at 6, 4.78% and 12.87% at 6-8 weeks of age, respectively (Table 4) This observation is in agreement with the finding of Kline et al. (1949). This may be due to the utilization of synthetic goitergens has produced variable increase in body weight and daily gain (Thrift et al., 1999). Weight loss with eltroxin than control and carbimazole by -22% and -24%, respectively at 8-10 weks of age. This weight loss is common in hyperthyroidism and caused by accelerated catabolism of feedstuffs and is accompanied by increased oxygen consumption. The loss in weight reflects a depletion of adipose tissue as well as muscle mass and often occurs despite an increase in caloric intake (Loeb, 1996b). Hypothyroidism effected indirect action by increasing in the receptors to increase protein synthesis and steroid hormones which are similar of GH and insulin response even though combined pre-parations an improvement in feed conversion efficiency and produced the best growth response (growth rate %), Buttery and Lindsay (1980). However, carbimazole enhanced body gain at 8-10 and 10-12 weeks than control and eltroxin by 61,86% and 55,24% at 8-10 weeks age, and 20.37% and 8.34% at 10-12 weeks of age, respectively. Also, it had improved growth rate and feed conversion than control by 8% and 3%, respectively during the whole experimental period.

Table 4: Means±SE of daily body gain (BG), growth rate (GR) and feed conversion (FC) of growing NZW rabbits in different experimental groups

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Items	Control	Eltroxin	Carbimazole
Average gain (g):			
5-6 weeks	17.6±1.8⁵	21.60±2.0 ^a	18.5±1.8 ^b
6-8 weeks	39.7±1.5 ^a	41.66±1.7 ^a	36.9±1.8 ^a
8-10 weeks	37.0±2.1°	38.58±2.2 ^b	59.9±1.5°
10-12 weeks	30.0±2.4 ^b	33.33±2.8 ^{ab}	36.1±2.5°
GR (%) (5-12 wk)	1.98±0.1	1.01±0.03	1.06±0.02
FC (5-12 wk)	4.1±0.1 ⁶	4.6±0.2 ^a	3.95±0.2 ^b

*. Values having different superscripts in the same row are significantly different (P<0.05). Carcass traits:

Carbimazole group had the higher final body weight than eltroxin and control groups by 11.1% and 13.5%, respectively (Table 5). This observation was agreement with the finding of Marai et al., (1994). The increase in body weight during the hypothyroidism is presumably a consequence of increased energy availability associated with a reduction in the basal metabolic rate (Thrift et al., 1999). Dressing percentage was significantly (P<0.05) higher in eltroxin and carbimazole groups than that of the control group. However, weights of the ovals including liver, kidneys, lungs and heart were almost higher in the control than both tested groups (Table 5). This observation for liver was agreement with the finding of Freake et al., (2001).

Blood hormones:

As affected by hypothyroidism, T_4 and T_3 concentrations significantly (P<0.05) lower as compared hyperthyroidism by -34.6% and -43.3%, respectively (Figs. 1 and 2). The hypothyroidism probably affects the secretion of T_4 directly and it could be the excessive conversion of T_4 to T_3 (Cooper et al., 1983). Hypothyroidism is known to decrease T_4 production from the thyroid gland and also to inhibit the extra thyroidal conversion of T_3 and T_4 (Achmadi and Terashima, 1995).

Table 5: Means ± SE of carcass traits in growing NZW rabbits in different

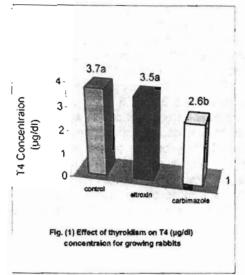
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Items	Control	Eltroxin	Carbimazole
Final body weight (g)	1850+61.24 ^b	1890+81.24ab	2100±68.92°
Dressing weight(%)	62.40±0.48b	64.25±0.32°	64.21±0.74°
Liver weight (%)	7.62+0.57	6.40±0.36	7.08±0.48
Kidneys weight (%)	1.49±0.08°	1.18 <u>+</u> 0.05 ^b	1.23±0.05 ^b
Lungs weight (%)	1.28+0.08	1.07±0.05	1.10 <u>+</u> 0.10
Heart weight (%)	0.60+0.03 ^a	0.45+0.41b	0.51±0.03°b

*. Values having different superscripts in the same row are significantly different (P<0.05).

The basal level of T_4 in control group increased (3.7±0.6 µg/dl) compared to hyperthyroidism (3.5±0.5 µg/dl) or hypothyroidism (2.6±0.3 µg/dl) (Fig. 1). Serum T_4 declined in hyperthyroidism implying that administration of exogenous T_3 caused group negative feedback at the level of the hypothalamus or pituitary to stop synthesis or secretion of thyroid stimulating hormone (TSH) (Thrift et al., 1999). Similar trend was observed for T_3 in control being (58.8±8.8 ng/dl) compared with (52.3±3.1) in hyperthyroidism and 36.5±6.3 in hypothyroidism (Fig. 2). The decline in T_3 concentration during the treatment period was reported by DeMoraes et al. (1998) for hypothyroidism. The decreasing in thyroid hormone in hypothyroidism being mor-nonounced in T_4 may be due to the process may reduce serum T_4 further, because T_3 production is still normal in spite of low absorption of follicular fluid, which results in low T_4 production. The organism probably compensates for the low metabolism following hypothyroidism by preventing the reduction of more metabolically potent hormone, T_3 (Shirpour et al., 2003).

Hematological parameters:

Results of hematological parameters in Table (6) show insignificant differences in red blood cell count (RBCs), white blood cell count (WBCs), platlet count, hemoglobin concentration (Hb) and packed cell volume (PCV), lymphocytes, monocytes, neutrophils distribution, mean corpuscular volume (MCV), mean corpuscular hemoglobin concentration (MCHC) and mean corpuscular hemoglobin (MCH) values. Similar findings were recorded by Rivlin and Wagner (1969), who found that anemia is uncommon in hyperthyroidism; however, in one study among 50 patients with hyperthyroidism 4 women were found with moderate anemia. Wintrobe et al. (1981) stated that administration of carbimazole is thought to be one of the least toxic of antithyroid drugs particularly in regard to its effects on lymph. Also, Peterson and Turrel, (1986) recorded that carbimazole is the drug of choice for long-term medical management of feline hyperthyroidism. In contrast to our results, Bond et al. (1983) observed some hematological changes including raised packed cell volume (PCV), erythrocytosis, occasionally mild anemia may be found in hyperthyroid cats. Differences between our results and those reported by pervious authors could be related to variety of aspects in the experimental design, such as dose of drugs, frequency, species, time that was determined to evaluate hematological changes.



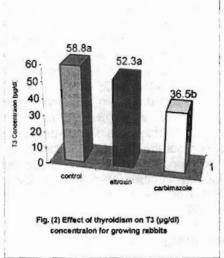


Table 6: Means <u>+</u> SE of hematological parameters of growing NZW rabbits in different groups

Items	Control	Eltroxin	Carbimazole
Red blood cells (x10 ⁶ /ml)	5.55±0.25	5.68±0.19	5.80±0.14
White blood cells (x10 ³ /ml)	7.02±1.24	5.53±1.03	5.24±0.25
Platlet counts (x10 ³ /ml)	324.20±82.05	236.00±50.51	223.40±30.43
Hemoglobin (g/100ml)	13.58±0.53	13.98±0.25	13,74±0.31
Hematocrit (PCV) (%)	38.34±1.72	38.28±0.91	38.46±0.89
Lymphocytes (%)	12.08±1.72	11.98±2.72	10.58±1.46
Monocytes (%)	14.88±1.63	15.87±2.53	14,84±0.73
Neutrophiles (%)	73.04±2.77	72.15±3.63	74.58±1.64
MCV*	68.40±0.68	67.00±1.06	66,80±0.58
MCHC*	35.74±0.34	37.15±0.95	36.14±0.38
MCH*	24.44±0.25	24.72±0.89	23.64±0.19

MCV mean corpuscular volume=PCV/RBCs

MCHC mean corpuscular hemoglobin concentration= Hb/ PCV

MCH mean corpuscular hemoglobin= Hb/RBCs

On the basis of the foregoing results, it could be concluded that the hypothyroidism improved growth performance for growing rabbits under summer condition in Egypt, reaching appropriate live body weight for marketing at shorter ages

REFERNCES

- Achmadi, J. and Y. Terashima (1995). The effect of propylthiouracyl-induced low thyroid function on secretion response and action of insulin in sheep. Domestic animal endocrinology 12:157-166.
- Amer M. A. and Chicago (1995). Physiology of thyroid hormones. IN: Division of drugs and toxicology, American Medical Association: Drug Evaluations Annual 1995, chapter 47, pp. 1039-1040.
- Baulieu E. and P. A. Kelly (1990). Hormones from molecules to disease. Hermann publishers in arts and science, chapman and hall, New York and London. Thyroid hormones pp. 368-374.
- Bond, B. R.; P.R., Fox and M. E. Peterson (1983). Echocardiographic evaluation of 30 cats with hyperthyroidism. Proc. Am. Coll. Vet. Int. med. 39.
- Bordy, S. C. (1945). Bioenergetics and Growth. Rein Hold Publishing Company, New York, USA. pp. 473-474.
- Burroughs, W., A. Raun, A. Trenkle and N. Raun (1960). Further observations upon the effects of methimazole upon feedlot performance and carcass characteristics of fattening beef cattle. Journal Animal Science 19:465-469.
- Buttery, P.J. and D. B. Lindsay (1980). Protein deposition in animals.

 Butterworths, London Boston, Sydney- Wellington- Durban Toronto.
- Cooper D.S., Kieffer J.D., Halpern R., Saxe V., Mover H., Maloof F., and Ridgway E.C., (1983). Propylthiouracyl (PTU) Pharmacology in the rat. II Effects of PTU on thyroid function. Endocrinology 113: 921.
- DeMoraes, G.V., H. R. Vera-Avila, A. W. Lewis, J. W. Koch, D. A. Neuendrorff, D. M. Haliford, J. J. Reeves and R. D. Randel (1998). Influence of hypo- and hyperthyroidism on ovarian function in brahman cows. Journal Animal Science 76:871-879.
- Dojana N.; A. Dinischiotu and M. Militaru (2000). The effect of thyroxine, insulin, hydrocortisone or adrenaline administration on pancreatic exocrine secretion in rabbit. Proc. 7th World Rabbit congress, Valencia, Spain. 4-7 July, 2000, vol. A pp:175-182.
- Duncan, D.B. (1955). Multiple range and multiple F test. Biometrics, 11: 1-42.
- Freake H. C., K. E.-Govon, K. Guda, C. Huang and S. A. Zinn (2001). Actions and interactions of thyroid hormone and zinc status in growing rats. The Journal of Nutrition vol. 131 No. 4 pp:1135-1138.
 - Greenspan, F. S. And J. D. Bsxter (1994). Basic and clinical endocrinology. App;eton and Lange Northwalk, Connecticut/San Mateo, California, 4th edition. pp:183

- Harada E., and S. Kato (1982). Influence of adrenaline, glucagons, hydrocortisone, thyroxine or insulin administration on pancreatic exocrine secretion in rats. Japanese Journal of Veterinary Science, 44(4): 589-596.
- Ingbar, S. H. and K. A. Woeber (1981). The thyroid gland of endocrinology, edited by R. H. Williams, 6th Edition pp 117-247. W. B. Saunders Company, Philadelphia, London, Toronto, Tokyo.
 - Iossa, S., L. Lionetti, M. P. Mollica, R. Crescenzo, A. Barletta and G. Liverini (2001). Fat balance and serum leptin concentration in normal, hypothyroid and hyperthyroid rats. *Int-j-obes-relat-metab-disord.*, 25(3):417-425.
 - Karsch, F. J., G. E. Dahl, T. M. Hachigian and L. A. Thun (1995) Involvement of thyroid hormones in seasonal reproduction. Journal Reprod. Fertil. Suppl. 49:409-422.
 - Kline, E. A., M. E. Ensminger, T. J. Cunha, W. W. Heinemann and W. E. Ham (1949). Effect of adding drugs to the ration of fattening cattle. J. Anim. Sci. 8:411-424.
 - LeGrow, A. B.; D. C. Fielding and T. A. Pressley (1999). Stimulation of Na, K-ATPase by hypothyroidism in the thyroid gland. Journal of endocrinology 160, 453-460.
- Loeb, J. N. (1996a). Metabolic changes in hypothyroidism. In: L. E. Braverman and R. D. Utiger (Ed.) The thyroid (7th Ed.), pp. 858-863. Lippincott-Raven, Philadelphia, PA.
- Loeb, J. N. (1996b). Metabolic changes in thyrotoxicosis. In: L. E. Braverman and R. D. Utiger (Ed.) The thyroid (7th Ed.). pp: 687-693. Lippincott-Raven, Philadelphia, PA.
- Marai, I. F. M.; K. A. El-Masry and A. S. Nasr (1994). Heat stress and its amelioration with nutritional, buffering, hormonal and physical techniques for New Zealand white rabbits maintained under hot summer conditions of Egypt. Proc. 1st Inter. Conf. on Rabbit production in Hot Climate, Cairo, Egypt, 6-8 September, 1994, pp. 475-487.
- Oyedipe, E. O., D. I. K. Osori, O. Akerejola and D. Saror (1982). Effect of level of nutrition on onset of puberty and conception rates of zebu heifers. Theriogenologh 18:525-539.
- Peterson, M. E. and J. M. Turrel (1986). Feline hyperthyroidism in : current veterinary therapy IX: small Animal prectico (ed. R. W. Kirk) pp:1026. W Bsaunders, Philadelphia.
- Rivlin, R. S. and H. N. Wagner (1969). Anemia in hyperthyroidism. Ann . intern. Med. 79:507
- SAS (1995), SAS User's Guide, Statistical Analysis System, Institute Cray, NC,USA.

- Shirpour A., S. Khameneh, V. Zarghami and M. Eskandari (2003). The influence of hypothermia on thyroid function in rats. Int. J. Endocrinol. 1: 27-32.
- Thoday k. L. (1993). Manual of small animal endocrinology. British Small Animal Veterinary Association, Kingsley House, Church Lan, Shurdington, Cheltenham. Gloucestershire Gl51 5TQ. Edited by Hutchison M. Spain, 2nd reprinted, pp. 41-57.
- Thrift T. A., A. Bernal, A. W. Lewis, D. A. Neuendorff, C. C. Willard and R. D. Randel (1999). Effect of induced hypothyroidism or hyperthyroidism on growth and reproductive performance of barhman heifers. Journal of Animal Science vol. 77 No. 7 pp:1833-1843.
- Tietiz N. W. (1995). Clinical guide to laboratory tests. 3rd ed. Philadelphia: W. B. Saunders (1995): 612.
- Wintrobe M. M.; G. R., Lee; D. R., Boggs; T.C., Bithell; J., Foerster; J. ., Athens and J. N., Lukens (1981). Clinical hematology eighth edition Lea and Fabiger, Philadelphia pp. 1038.
 - تأثير مستوى نشاط الغدة الدرقية على الأداء الإنتاجي في أرانب النيوزيلندي الأبيض النامية تحت ظروف الصيف الحار في مصر
 - محمد عبد العزيز الصاوي * / هدي عبد الرعوف شعبان " وعبد الفتاح عبد الحميد ندا " " معهد بحوث الإنتاج الحيواني مركز البحوث الزراعية وزارة الزراعة ـ مصر.
 - *معهد بحوث صحة الحيوان مركز البحوث الزراعية وزارة الزراعة ـ مصر.
 - أستخدم في هذه الدراسة عدد ٣٠ أرنب نيوزيلندي أبسيض عمسر ٣٥ يسوم بمتوسسط وزن ٢٦ ±٢٠ جم وقد تم تقسيمهم إلي ثلاثة مجاميع متشابهة عوملت بالحقن تحت الجلسد مسرة واحسدة أسبوعيا لمدة ٧ أسابيع كالتالى:
 - مجمُّوعة أولى : (مجمُّوعة المُقارنة) تم حقنها ٥٠٠مل محلول ملح فسيولوجي.
 - مجموعة ثانية : تم حقنها بالالتروكسين (لرفع نشاط الغدة الدرقية) ١٥ ميكروجرام/ارنب .
 - مجموعة ثالثة : تم حقنها بالكاربيمازول (لخفَّض نشاط الغدة الدرقية) ١٥ ميكروجر لم/ارنب .
 - أوضحت النتائج زيادة كمية الغداء الماكول ووزن الجسم والتحسين في معدل النمو والاستفادة الغذائية لمعاملة الكاربيمازول عن الألتروكسين والكنترول ولكن لوحظ انخفاض درجة حرارة المستقيم للمعاملة بالألتروكسين في عمر ١٢ أسبوع. أدت المعاملة بالإكاربيمازول زيادة الوزن النهائي عسن مجموعتي الألتروكسين والكنترول بمعدل ١٠١١% و ١٣٠٥% على التوالي.
 - وكذلك وجد زيادة معنوية على مستوى ٥% في نمية النصافي لمجموعتي الالتروكسين والكاربيمازول مقارنة بمجموعة الكنترول على الرغم من زيادة أوزان كلا من الكبد والكليتين والرنتين والقلب فيها عن المجموعتين الأجرتين المعاملتين. وقد لوحظ أن ارتفاع مستوي هرموني الثيروكسين والتراي أيودوثيرونين في مجموعة الكنترول عن المجموعتين المعاملتين، ولم يكن هناك أي تأثيرات معنوية للمعاملات على الخصائص الهيماتولوجية للدم .