# EFFECT OF ADMINISTRATION OF L-TYROSINE AND L-CARNITINE ON ONSET OF PUBERTY IN RABBITS

NOSEIR W.M.B., ABBOUD M.Y.A., AMRAWI G.A., LOTFY T.M.E. Faculty of Veterinary Medicine, Alexandria Uni.

Received: 28. 11. 2007 Accepted: 10. 12. 2007

### **SUMMARY**

The present study was conducted on 72 New Zealand white rabbits assigned into two age groups (two and three months): 36 rabbits for each group of the same age (18 males and 18 females). For. treatment, 24 rabbits (males and females) were given orally a single dose of L-tyrosine, 100 mg/ kg body weight; the other 24 rabbits (males and females) were given a daily oral dose of 1 gm Lcarnitine/kg. The other 24 rabbits were left as control. All rabbits were kept under complete observation during the experimental period and data obtained were statistically analyzed. There was a steady increase in the body weight with advancement of age; it seemed to be similar for males and females. The treated group revealed a significant higher body weight (P<0.05) when compared to control and treated females, exhibited higher body

gain than males. In the treated group, the testicular descent in males and the first onset of male acceptance by females occurred much earlier during the second month of age compared to control. Successful mounting and ejaculation of semen containing spermatozoa for the first time was recorded in the treated bucks at the beginning of the third month of age, much earlier than in control. Testes and ovaries size increased significantly with advancement of age. L-tyrosine and Lcarnitine treated groups revealed significant increase in the testicular and ovarian size compared to those in the control group (P<0.01). It could be concluded that L-tyrosine and L-carnitine, both act as growth promoting factors. They could be used in rabbits for enhancement of puberty in both males and females.

### INTRODUCTION

For several reasons, rabbits are unique among small animals for food and commerce. They produce highly nutritious, low-fat and lowcholesterol meat which is rich in proteins, certain vitamins and minerals. Being herbivores, they do not compete with humans for their food and are easily adaptable to different environments, and investment and labor costs are low. They are easy to transport and market for food, fur and skin. In addition they are highly productive, characterized by long breeding season, which may extend from October to May, associated with a large number of litters with short gestation and lactation periods (Daader et.al., 2002). Puberty of the male rabbit has been defined as the first time when the male becomes capable for producing sper-matozoa in the ejaculate (Sand, 1979). It has also been defined as the stage when the endocrine function ofthe testes first becomes clearly evident (Donovan and Van der werfften Bosch, 1965). Puberty occurs a month or even more before rabbits attain sexual maturity, and there in an age difference between the different breeds and strains of rabbits (Sand, 1979). The small white Polish bucks reach puberty at four months of age, whereas New Zealand white and Californian bucks reach puberty at five months of age (Peter, et.al., 1979).

Based on the use of some organic substances used to improve the reproductive efficiency of animals through enhancing the release of sex hormones,

the present study aimed at studying two of these substances: L-Tyrosine and L-Carnitine. L-Tyrosine is a semi essential amino acid involved in the formation of catecholamine from adrenal gland and thyroxin in the thyroid gland (Harmmerl and R,sse, 1987). L-Carnitine is a dipeptid amino acid. Sex hormones are steroid hormones, and all steroid hormones are ultimately made from a single precursor, cholesterol, which in turn is made from acetyl-CoA that is formed through the carnitine transport (Lehninger, 1982).

j.

0

M

N.

135

(of

99.

18

f ti

ani

 $\Lambda^{(1)}$ 

z)K

jelpl

 $\Delta U^{2}$ 

to et

[.01

MAT

**Inim** 

mma!

nonth

Medic

od fro

lonsed

22555

ad f

1000

y mir

ब्बहुed र्षः of

have

<sup>V</sup>etMe

L-Carnitine performs the following tasks in relation to spermatogenesis and function of mammalian spermatozoa; 1) Synthesis and protection of cell membranes (Uhlenbruck, 1996), 2) Storage of activated acetic acid in the form of acetylcarnitine, which is an energy-rich compound that serves as the primary energy source for sperm motility after insemination (Jeulin et.al., 1988). Trials have indicated that, L-carnitine has beneficial effects in both productive and reproductive performance of monogastric species including sows (Harmeyers, 1993 and Freemaut et. al., 1993), mares (Iben and Leibertseder, 1994) and poultry (Leibertseder and Iben, 1994). In contrast to monogastric species the energy supply of ruminants is based almost entirely on oxidation of short and long chain fatty acids rather than carbohydrates. L-carnitine is therefore, even more important for ruminant than monogastric species, Noseir and Amrawi (2000) found an improvement effect of L-carnitine supplementation on fertility

Vet.Med.J., Giza. Vol. 55, No. 4(2007)

1112

of normal and sub fertile rams.

L-tyrosine has been found to act as a growth promoting factor and hormonal inducing factor in rabbits (Abo-Elroos, 1992), rats (Scheuermann, 1984), dogs (Aumann, 1987), pigs (Gindele and Koppen, 1988), and buffaloes (Amrawi et.al., 1992; Amrawi, 1997). It is of interest to study the changes in activities of gonads by administration of those amino acids, however, it is difficult to confirm these changes in mature animals in which gonads are normally functioning, and therefore selecting animal just before reaching puberty is helpful to study the effect of these amino acids on gonads. The present study was designed to study the effect of administration of either L-tyrosine or L-carnitine on the onset of puberty in rabbits.

## MATERIALS AND METHODS

Animals: Seventy two (36 males and 36 females) immature New Zealand white rabbits aged 2-3 months and bred at the Faculty of Veterinary Medicine, Alexandria University during the period from January to April were used. Rabbits were housed individually inside steel cages, with free access to clean water via automatic water troughs and feeding on a constant plan of nutrition throughout the period of the experiment. The daily minimum and maximum indoor temperatures ranged between 14 and 20°C. A lightening schedule of 12h of light was used. Each animal was allowed a daily non-treated ra-tion composed of

about 200g concentrates (including 27% barley, 25% wheat bran, 20% ground corn, 15% ground horse beans, 8% soybean meal, 1.5% bone meal, 3% minerals and vitamins, and 0.5% common salt), in addition to barseem hay ad libitum. Rabbits were given a period of adjustment for at least two weeks before reaching the starting age of experiment (2 and 3 months), to ensure a good health condition and to avoid stress during transportation, through the daily observation of any clinical signs indicating infection or parasitic infestation.

Experiment: Rabbits were assigned according to the age at the beginning of the experiment into two equal groups; A and B aged 2 and 3 months, respectively. Each group contained, 18 males and 18 females;

Group A (2 months old): both sexes were divided into 3 equal subgroups: the first of which (6 males and 6 females) was treated by administering 100 mg L-tyrosine/kg body weight (Theriogon-ADWIA), in a single oral dose by direct dilution into 3 ml sterile water. Each animal received a separate dose orally. The second subgroup (6 males and 6 females) was treated by oral administration of 1gm L-carnitine/kg body weight (Carniking 50%-Pharopharma Egypt) by direct dilution into 5 ml sterile water. Each animal received the dose as in the first subgroup and the treatment was given daily for two weeks. The last subgroup (6 males and 6 females) was kept as control.

of age, while in L-carnitine treated group the testicular descent occurred at 14-15th day after the second month of age. This was remarkably earlier than in the control (27-28th day). When females were checked daily for standing to be mated by males as a sign of sexual development, it was noticed that the first time for male acceptance by females was much earlier in tyrosine-treated group (10-11th day) and in carnitine group (13-15th day) beginning from the second month of age compared to control (21-23rd day).

4- Sexual interest was recorded in the tyrosinetreated bucks at the beginning of the third month of age (90-97 days), while in carnitine group at 85-95 days, whereas control bucks displayed sexual interest at approximately the middle of the fourth month for the four months old rabbits (131-133 days).

Changes in the testicular and ovarian size of experimental groups are presented in Table, 3. There was a significant statistical difference (P<0.01) in ovarian size between tyrosine treated group and both control and carnitine groups.

Number of kids in treatment groups (Table, 4) in both experimental groups (A&B) were significantly (P<0.05) higher than control.

Table (1):- Effect of age and treatments (L-tyrosine and L-carnitine) on body temperature of rabbits (°C)

n ja	Group A (2 Months old)	Group B (3 Months old)	Overall mean
L-Tyrosine	38.45±0.34 a	38.59±0.15 a	38.51±0.22 a
L-carnitine	38.55±0.08 °	38.73±0.11 a	38.61±0.10 a
Control	38.31±0.24 a	38.48±0.17 a	38.41±0.19 a

Means  $\pm$  SE. Means with different alphabetical superscripts (a,b,..) in the same column are significantly different (P<0.05).

Table (2):- Effect of age, sex and treatment (L-tyrosine and L-carnitine) on body weight and net body gain of rabbits.

Group (A)	Starting b.w.(kg)	After 1 Month(kg)	Net Weight gain(kg)
Male	le		
L-tyrosine	2.16±0.25	2.89±0.16	0.73±0.29 a
L-carnitine	2.00±0.31	2.71±0.17	0.71±0.31 a
Control	2.43±0.06	2.85±0.15	0.42±0.21 b
Female :			
L-tyrosine	2.05±0.42	3.12±0.17	1.07±0.16 a
L-carnitine	2.00±0.44	3.06±0.55	1.06±0.27 a
Control	2.33±0.11	3.15±0.38	0.82±0.23 b
Group (B)	Starting b.w. (kg)	After 1 month(kg)	Net Weight gain(kg)
Male			Chillian Charles
L-tyrosipe « .	2.80±0.08	3.10±0.12	0.30±0.10 a
L-carnitine	2.86±0.12	3.12±0.17	0.26±0.08 a
Control	2.80±0.04	2.99±0.04	0.19±0.09 b
Female			L J = 11, 12
L-tyrosine	3.11±0.56	3.43±0.77	0.32±0.06 a
L-carnitine	3.17±0.18	3.49±0.39	0.32±0.08'a'
Control	3.20±0.40	3.52±0.61	0.32±0.04 a

Means  $\pm$  SE. Means with different alphabetical superscripts (a,b,..) in the same column are significantly different (P<0.05).

b.w. = body weight (kg).

Vet.Med.J., Giza. Vol. 55, No. 4(2007)

each of the most flated case please missisfers it is not a

3 1

1116

Table (3):- Interaction of age and treatment (L-tyrosine and L-carnitine) on Testicular and ovarian size of rabbits (cm<sup>3</sup>).

	Group A (2 Months old)	Group B (3 Months old)	
Testicular size			
L-carnitine	2.59±0.05 *	2.85±0.20 *	
L-tyrosine	2.61±0.04 a	3.06±0.10 <sup>a</sup>	
Control	1.59±0.10 b	1.82±0.06 b	
Ovarian size	mg <sup>T</sup> to 1	, "	
L-carnitine	1.03±0.08 a	1.09±0.03 <sup>a</sup>	
L-tyrosine	1.24±0.07 a	1.41±0.07 b	
Control	0.46±0.04 b	1.06±0.04 a	

Means ± SE. Means with different alphabetical superscripts (a,b,..') in the same column are significantly different at P<0.01.

Table (4):- Interaction of age and treatments on the number of kids.

Laxui seri	Group (A)	Group (B) ·	Overall mean
L-carnitine	6.5±1.2 a	6.7±0.9 a	6.6±1.0 a
L-tyrosine	6.7±1.5 a	6.5±1.2 a	6.6±1.3 a
Control	4.7±0.9 b	5.5±1.0 b	5.1±0.9 b

Means  $\pm$  SE. Means with different alphabetical superscripts (a,b,..) in the same *column* are significantly different at P<0.05.

# DISCUSSION:

It is well known that a phenomenon of the body metabolism interpreting a change in the physiological activity of an in-dividual is to find a corresponding change in the body tempera-ture. Thus, increasing the body metabolism seems to have a posi-tive relationship with body temperature. Although, in the present study, differences in the body temperature recorded (Table, 1) among experimental and treatment groups were not significant, but higher body temperature ap-peared with the rabbit approaching age of puberty was observed in both treatment groups.

In the meantime advancement of age of rabbits under the present experimental condition was noticed to have a significant positive relationship to the body weight. The net gain in the body weight appeared more prominent in rabbits of two and three months (Table, 2), a finding which also supported the previous suggestion that puberty in rabbits starts with the third month of age (Lebas et.al., 1986), contrary to that mentioned in some earlier studies (Arrington and Kelly, 1976; Peter et al., 1979; Sand, 1979). Throughout the different ages when tyrosine was administered, the body weight of the treated rabbits revealed a highly significant increase when compared to that of control. It could be speculated that tyrosine acts as a growth promoting factor. This is in accordance with the earlier reports showing that increasing the circulating tyrosine activates the hypothalamic catecholamine, dopamine, which in turn activates the release of growth hormone (M,ller, 1973), in addition to its importance in formation of thyroxin and protein synthesis (Harper et al., 1980).

Throughout the age of two months group when carnitine was supplemented, it produced positive and significant effect on growth rate compared to control. This finding comes in accordance with studies conducted on sows (Musser et. al. 1999).

From the present study, it has been shown that testicular descent into the scrotum takes place much earlier during the second month of age in the treated groups compared to control. Klaws (1985) and Lebas et al., (1986) indicated that the testicular descent into the scrotum of rabbits takes place shortly before puberty at the age of 3-4 months. This could be recognized as a sign for the steroid ac-tivity of the testis in the male before reaching the expected age of puberty in rabbits. It has long been known that the testicular descent is completed under the influence of androgens secreted by the testes (Hafez, 1986).

Carnitine has an important role in spermatogenesis regarding synthesis and protection of cell membranes (Uhlenbruck, 1996), beside storage of activated acetic acid in form of acetylcarnitine, which is an energy-rich compound that serves as

Vet.Med.J., Giza. Vol. 55, No. 4(2007)

1118

the primary energy source for sperm motility after insemination (Jeulin et.al., 1988). On other hand, L-tyrosine might induce early puberty in male rabbits as a growth promoting factor (M,ller, 1973) and a stimulant to the gonadotrophin releasing factor (Kamberi et. al., 1971) which potentiates the testicular function. The sexual behavior of females is synchronized by the ovarian activity (Hafez, 1986), as the presence of developing follicles in the ovaries is followed by secretion of estrogen resulting in the female standing position to be mated by the male. This support the hypothesis that tyrosine functions in the synthesis of cerebral catecholamine, this was shown to activate the hypothalamus to release GnRH (Hammerl and R,sse, 1987; Arthur, 1989) which stimulates the Pituitary gland to secrete gonadotrophins (FSH & LH) activating the ovaries. Kamberi et al., (1971) reported that dopamine, created from the amino acid precursor L-tyrosine, might function as a synaptic transmitter to potentiate FSH releasing factor or LH-Releasing factor or both.

took speci

ed posting

on pared to

tace with

1,1999)

town to

tes pla

of apr

ol. No.

deate

bis tals

104

to for the

k kin

zióis t

exel 8

2 60%

MONCH.

of of

or!

13

no f

It is of interest to notice that the testicular size of rabbits throughout the experimental period appeared remarkably larger in the tyrosine as well as carnitine treated groups compared to control in both ages (A and B), a finding which confirms the close relationship between the testis weight and body weight, and also the relation between the testis weight and the age of the animal (Amann, 1970). Similarly, the ovarian size showed a highly

significant increase in the tyrosine and carnitine treated rabbits when compared to the control group (Table, 3). L-carnitine was highly effective in rabbits at two and three months of age where the ovarian weight in the treated cases was higher than that of the control. On contrast with tyrosine treated females, where the ovarian size was remarkably high in ages, two and three month groups than both carnitine and control groups. Thus, if it has been supposed that puberty in rabbits takes place at four months age (Harkness and Wagner, 1989), treatment with tyrosine and carnitine promotes the developing process of gonads to module puberty at a stage earlier than that under the natural conditions.

When comparing the treatment effects on the number of birth, it reveals that in treatments groups, tyrosine and carnitine, the numbers of kids were higher than that of control in both ages. These results however differ than that of Musser et.al. (1999) on pregnant sows where, they postulated that carnitine does not affect number of birth rather than litter weights and viability.

From the clinical observations and results of the present study it could be concluded that both amino acids L-tyrosine and L-carnitine could have the following physiological functions in rabbits;

1) Growth promoting factor indicated by the prominent increase in body weight and sizes of gonads. 2) Enhancement of puberty in both males

Vet.Med.J., Giza. Vol. 55, No. 4(2007)

**CS** CamScanner

and females.

# REFERENCES

Abo-Elroos, M.A. (1992): Effect of tyrosine on the pituitary and gonads of rabbits in relation to sexual maturity. M.Vsc. Thesis, Zagazig Uni., Moshtoher-Benha Branch. Amann, R.P. (1970): The male rabbit, quantitative testicular histology and com-parisons between daily sperm production as determined his-tologically and daily sperm output. Fertility and Sterility, 12:662-672.

Amrawi, G.A. (1997): The Effect of L-tyrosine on fertility of bull and buffalo-bull. Alex.J.Vet.Sci.13: 45-54.

Amrawi, G.A., Hussein, F.M., El-Bawab, Iman B. and Zakaria, A.D. (1992): The Induction of puberty in delayed pubertal buffalo- heifers by L-Tyrosine. Cairo International Meeting on Working Animals. 12-14 April, NIR, Cairo, Egypt.

Arrington, L.R. and Kelly, K.C. (1976): Domestic Rabbit biology and production. Gainesville, The University Press of Florida.

Arthur, R.J. (1989): Veterinary Reproduction and obstetrics. 6th Edition, Bailliere Tindall, London.

Aumann, J. (1987): Beitrag zur Behandlung von fertilitatsstorungen der deutschen schaferhudin. Diss. Med. Vet., M.nchen.

Daader A.H., Gabr, H.A, Khadr, A.M.F., and Seleem T.S.T. (2002): Fertility traits in different breeds of rabbit does as affected by coitus frequency and remating interval.

3rd Scientific Conference on Rabbit Production in Hot Climates. 8-11 Oct., Hurghada, Egypt.

Donovan, B.T. and Van der Werfften Bosch, J.J. (1965); Physiology of puberty. Edward Arnold, London. Freemaut, G., Letre, J. and Aerts, J. (1993): Do lactating sows benefit from L-carnitine supplementation. Varkensbedrijf (june): 30.

Gindele, H.R. and Koppen, W.D. (1988): Can the reproductive performance of sows be improved by oral administration of L-tyrosine. Tierärzt. Umschau, 44:528-534.

Hafez, E.S.E., (1986): Reproduction in Farm Animals. Laboratory animals, 5th Ed. Lea and Febiger.

Hammerl, J. and Russe, M. (1987): Der einfluss der aminosaure L-tyrosin und L-tryptophan auf das fruchtbarkeitsgeschehen bei jungen Ratten. Zuchthygiene, 22:80-84.

Harkness, J.E. and Wagner, J.E. (1989): The Biology and Medicine of Rabbits and Rodents, 3<sup>rd</sup> Edition, Lea and Febiger, Philadelphia.

Harmeyers, J. (1993): The effect of additional L-carnitine at the end of gestation and during lactation on sow and litter performance. Internal. report for Lonza Ltd., M, nchensteinstrasse 38, C4002 Basel.

Harper, H.A., Rod-well, V.W., and Mayes (1980). Review of physiological chemistry 17<sup>th</sup> edition. Long Medical publications, Middle East Edition PP: 265-272.

Iben, C. and Leidbertseder, J. (1994): Camitine in Horse. Proceeding of the 4th annual congress of the European society of Veterinary Internal Medicine (ESVIM), Brussels, (Belgium) pp19.

Ismail, S.T. (1979): Factors affecting sperm production in Camel. M.V.Sc. Thesis, Cairo Uni.

Jeulin, C., Soufir, J.C., Marson, J., Paqulgnon, M. and Dacheux, J.L. (1988): Acetylcarnitine and spermatozoa: relationship between epididymal maturation and motility

- in boars and man. Reprod. Nutr. Dev. 28:1317-28.
- Kamberi, I.A, Mical, R.S. and Porter, J.C. (1971): Effect of, anterior pituitary infusion and intraventricullar injection of catecholamine on prolactin release. Endocr. 88:1012-1020.
- Klaws. J.L. (1985):A compendium of rabbit production. Agriculture and Rural development, Eschborn.
- Lebas, F; Coudert, P; Rouvier, R and Rochambaa, H. (1986): The Rabbit, husbandry, health and production. FAO Animal production and health series No. 21.
- Lehninger, A.L. (1982): "Principles of Biochemistry" CBS Pub. and Distrib., Shahdra, Delhi, India.

-

Leidbertseder, J., and Iben, C. (1994): Carnitine in poultry.

Proc. of the 4th annual congress of the European Society of Veterinary Internal Medicine (ESVIM), Brussels, (Belgium) pp17.

11

- Müller, E.E. (1973): Nervous control of growth hormone secretion. Neuroendocr. 11:338-369.
- Musser R.E., Goodband R.D., Tokach M.D., Owen K.Q., Nelssen J.L., Blum S.A., Dritz S.S. and Civis C.A. (1999): Effects of L-carnitine fed during gestation and lactation on sow and litter performance. J. Anim. Sci. 77: 3289-95

E.

W 4

A 170

D

- Norusis, M.J. (1986): SPSS/PC+for the IBM PC/XT/AT. SPSS Inc., Chicago.
- Noseir, W.M.B. and Amrawi, G.A. (2000): L-carnitine supplementation in normal and subfertile rams.12th Annual Congress, Egyptian Soc.Anim. Reprod. Fert., pp137.
- Peter, R., Nephi, M.P. and George, S. (1979): Rabbit production. The Interstate Printers and Publishers, Danville, Illinois.
- Sand, J.C. (1979): The Domestic Rabbit. 3rd. Ed. John Wiley and Sons., New York.
- Scheuermann, R. (1984): Zur bedeutung der aminosäure tyrosin im reproduktion sgeschehen der ratte. Diss. Med. Vet. M,nchen.
- Uhlenbruck, G. (1996): L-carnitine and the immune system: from the mode of metabolism to the modulation of membranes. In: Carnitine-Pathochemical Basics and Clinical Applications. Eds Selm, H., Loster, H. Ponte press Bochum. pp 47-60.