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SERUM AND SEMINAL THYROID HORMONES IN RELATION TO SEMEN QUALITY OF FRIESIAN BULLS

BY

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INTRODUCTION

T4 and T3 are iodoamino acid hormones produced by the thyroid gland. The biological halflife (t½) of T4 is 4-5 times that of T3 as well as T3 binds to the receptor of target cells with 10 times as the affinity of T4 (Schimmel and Utiger, 1977 and Martin et al., 1985). Moreover, the majority of circulating T3 arises by the peripheral monodeiodnation of T4 in the liver and kidneys (Sterling et al., 1977).

In farm and laboratory animals, hypothyroidism or thyroidectomy resulted in degeneration of the testis, arrest of spermatogenesis as well as depression of spontaneous activity, motility, viability and increased abnormality of spermatozoa (Swanson and Boatman, 1953, Bruni et al., 1975 and Ghorieb et al., 1978). On the other hand, thyroid therapy was suggested to improve reproductive performance of the male animals (El-Azab et al., 1974).

The metabolic energy required for survival and motility of spermatozoa is mostly gained from seminal fructose through fructolysis. Moreover, an intimate relationship was found between fructolytic rate and spermatozoal density and motility in bull semen (Mann and Lutwak-Mann, 1981).

Available literature concerning existance and content of T₄ and T₃ in semen of bulls are scarce. Therefore, this study was planned to determine levels of serum and seminal T₄ and T₄ as well as their relation with semen quality.

MATERIALS AND METHODS

Sixteen mature, healthy, fertile Friesian bulls subjected for artificial insemination puropse in Beni Suef Centre for Artificial insemination were used. The age of bulls ranged from 3-5 years and their mean body weight was 500-600 kg. All bulls were under veterinary supervision and tested periodically against tuberculosis, brucellosis trichomoniasis and vibriosis. They were fed on barseem (30 kg/bull daily) and a concentrated ration (6 kg/bull daily) composed of 65% cotton seed cake, 20% wheat bran, 12% rice polish, 2% lime salt and 1% common salt. For this study, periodical semen samples were collected from these bulls by using artificial vagina according to Walton technique (1945) for 4 successive weeks to evaluate their semen quality (sperm density, motility %, live % and abnormalities %). Accordingly, bulls were divided into 2 groups; the first (10 bulls) was of high semen quality, while the second (6 bulls) was of moderate quality. Furthermore, individual semen samples were collected from both groups and represented the first and second ejaculates. As soon as semen sample was obtained, it was divided into 3 parts; the first was used for evaluation of

semen characters (Salisbury and Van-Demark (1961). The second part was centrifuged directly at 3000 rpm for 20 minutes in order to separate seminal plasma which was kept at -20°C till estimation of Thand Ta contents by 1125 radioimmunoassay according to Abraham (1981). T4 and T3 radioimmunoassay kits were supplied by ICN Biochemical Inc, Diagnostic Division. The third seminal part was used to determine fructose concentration at zero time as well as after 1, 2 and 3 hours incubation at 37°C and the fructolysis index was calculated according to Mann (1948). In the same time, individual blood samples were collected from these bulls early in the morning. The sera were separated and preserved at -20°C till radioimmunoassay of T4 and T3 as performed for seminal plasma.

Statistical analysis of the obtained data and correlation factor "r" were carried out according to Snedecor and Cochran (1967).

RESULTS

As demonstrated in Table (1), bulls of high semen quality showed a significantly higher (P < 0.01) sperm concentration, motility % and live % as well as lower abnormalities % (1 st ejaculate) than their corresponding values of bulls with moderate semen quality.

Data of Table (2), clarified that serum T_4 and T_3 levels were significantly higher (P < 0.02) in bulls of group I than those of group II. Aslo, T_3 contents in the seminal plasma of bulls in group I were higher than their corresponding values in bulls of group II. Within each group, T_3 content was higher in the 1 st ejaculate than that of the 2 nd one. On the contrary, T_4 seminal levels of bulls in group I were lower than

Table 1 : Semen characteristics of Friesian bulls with high and moderate semen quality. 13.48 ± 1.438 Abnormalities 11.21 ± 0.97 10.32 + 0.75 72.96 ± 2.45 b 73.64 ± 2.018 84.98 ± 2.94 b 85.37 ± 3.25ª 63.97 ± 1.54 b 83.41 ± 1.368 84.22 ± 1.23 b 60.97 ± 1.13⁸ Motility concentration 0.73 ± 0.04 b 1.06 ± 0.048 1.73 ± 0.05 b 0.82 ± 0.03ª Sperm cell (X 109 1 st ejeculate 1 st ejaculate 2 nd ejaculate 2 nd ejaculate II. Bulls of moderate semen quality semen quality I. Bulls of high Groups

In the same column, values having the same letters differ significantly at P 6 0.01.

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Table 2 : Serum and seminal T4 and T3 concentrations as well as fructolysis index in Friesian bulls.

		Serum		ed a rg b	(Semen	30
Groups	T4 (ag S)	T ₄ T ₃ (ng S)	T4/T3 retio	T4 (418 %)	T ₄ T ₃ (11g K)	T ₄ /T ₃ retio	Fructose concentration Fructolysis index at zero time (mg/109 sperm/hou et 37°C)	Fructolysis index (mg/109 sperm/hour at 37°C)
I. Bulls of high semen quelity	15.81±	15.81± 380.72± 0.28 ⁸ 5.99 ⁸	41.53	#4VsT#71# 11@438## 40darn188	rani io n	go podkiol godenika elemenos		
1 st ejeculate	district			0.85±	0.85± 276.95± 0.02 ⁸ 5.48 ⁸	3.07	261.35± 10.38 ⁸	1.10±
2 nd ejeculete	fand.			1.34±	1.34 166.38± 0.088 8.658	8.05	426.10± 17.67ªbc	1.49±
II. Bulls of moderate 14.07± 328.27± semen quality 0.418 4.708	14.07±	328.27±	42.86	# 17	na le Le la			
1 st ejeculate	13 00 1000 79 yl			13.24±	13.24± 257.34± 0.54 ⁸ 4.48 ⁸	51.45	245.15± 14.76 ^b	0.70+
2 nd ejeculete	70,000 120518 120518	820e-13 202-11		16.05±	16.05± 125.67± 0.37 ⁸ 5.35 ⁸	127.71	e Seedall	0.93±

In the same column, values have the same letters differ significantly at P 4 0.02.

those of group II. The lst ejaculate, within each group, contained significantly lower values of T4 than the 2 nd ejaculate. The fructolysis index of the first group was greater than that of the second group. Then within each group, the 2 nd ejaculate had a higher index than that of the lst ejaculate.

DISCUSSION

Tables 1 and 2 revealed that bulls with high semen quality (group I) possessed higher serum T4 and T3 levels than those having moderate quality (group II). This finding indicates the intimate relationship between serum T4 and T3 levels and male reproduction and supports previous studies of Goswami (1964); Soliman et al. (1978) and Soliman and E1-Toukhy (1980) who reported that administration of thyroid hormones increased the activity of spermatogenesis and spermiogenesis. In this respect, the influence of thyroid hormones could be attributed to their effect upon synthesis, release and turnover of pituitary gonadotropic hormones (Bruni et al., 1975; Soliman et al., 1978 and E1-Toukhy et al., 1982).

Previous studies regarding presence of T4 and/or T3 in semen and their interaction with semen quality are scarce and do not give a meaningful conclusion. The present findings showed that sperm density was directly proportional with seminal T3 content (r = 0.943) and inversely related with seminal T4 (r = -0.913). Thus, semen samples with the highest sperm count (group I, lst ejaculate) contained the highest T3 and the lowest T4 contents. This could be explained by the findings of Schimmel and Utiger (1977) and Sterling et al. (1977) who reported that 65-75% of the metabolic effect of thyroid hormones is due to T3; also 33-40% of circulating T4 is monodeiodinated into T3. Moreover, Armstorous et al. (1982)

reported that intravenous injection of T₄ into bulls resulted in an increased seminal T₄ and T₃ contents after 2 hours.

The present data cleared out that serum T4/T3 ratio was nearly similar in both groups (I and II). On the other hand, seminal T4/T3 ration exhibited exaggerated higher values in gorup II than in group I. Also, a reverse relationship was found between semen quality and seminal T4/T3 ratio. This observation motivated the hypothesis that inhibited conversion of T4 into T3 leads to accumulation of T4 in semen and creates an imbalance in T4/T3 ratio which may be responsible for decreased seminal quality of bulls in group II. Therefore, it is advisable to include the determination of seminal T4/T3 ratio during the judgment of semen quality in bulls.

Concerning fructolysis index, Table, 2 showed that in group I it was sugnificantly higher (P < 0.02) than that of group II. Besides, the index was inversely related with seminal T4/T3 ratio in both groups. The observed increase of fructolysis index in the 2nd ejaculate than the 1st, in each group, may be attributed to the higher concentration of fructose at zero time in these ejaculates. Amir et al al. (1965) reported that there was a highly positive correlation among fructolysis, semen quality and initial fructose concentration.

In this study, the obtained low T4/T3 ratio in bulls with high semen quality reflects the importance of existance of metabolically active T3 hormone in semen. This hormone may play an important role in energy production through fructolysis. Martin et al.(1985) reported that thyroid hormones increase the cyclic adenosine monophosphate (cAMP) concentration through enhancing the synthesis of adenylate cyclase. Moreover, Casillas and Hoskins (1970 and 1971) reported

that cAMP synthesising system in mammalian spermatozoa is insensitive to a wide variety of hormones
except 3, 3, 5 triiodo-L-thyronine (T3) which speaded up the synthesis of cAMP. Mann and Lutwak-Mann
(1981) reported that generation of cAMP from ATP by
adenylate cyclase has a stimulatory action on motility of spermatozoa. This motility includes the
passage of spermatozoa from caput to cauda epididymidis, motility at ejaculation time and motility in
the female genital tract during capacitation.

Thus, it is essential for bulls, especially those used for reproductive purposes, to feed on a ration containes balanced amounts of building stones of thyroid hormones, besides periodical evaluation of serum and seminal T₄/T₃ ratio to avoid disturabnces in thyroid hormones level which is considered one of the factors leading to subfertility in bulls.

SUMMARY

Serum and seminal levels of thyroxine (T4) and triiodothyronine (T3) were evaluated and correlated with the semen quality of two groups of Friesian bulls. The first group (I) was of high semen quality while the second (II) represented bulls of moderate quality. Obtained data clarified that serum T4 and T3 levels were significantly higher in bulls of group I than those of group II. Serum T4/T3 ratio was nearly similar in both groups. Seminal plasma T3 contents of group I were higher than their corresponding values of group II, while T4 concentrations of the second group were sharply greater than those of the first group. It was concluded that a reverse relationship is found between seminal T4/T3 ratio and semen quality. T3 has the upper hand in regula ting semen activity. Decreased conversion of T4

into T₃ in semen leads to an imbalance among both hormones which could be considered as a factor responsible for inhibited sperm activity.

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