The Effects of Trenbolone Supplementation on the Extremity Bones in Running Rats

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Abstract

Anabolic steroids are testosterone derivatives through which anabolic effects are maintained and androgenic effects are minimized. The use of ergogenic agents is increasing among athletes for doping in order to increase physical performance and change external image. The objective of this study was to determine effects of trenbolone supplement administered on running rats for 4 weeks on extremity bones. The study was conducted with 28 male Wistar rats aged 28 days with a mean weight of 61.80 g supplied from the Selcuk University Experimental Medical Research and Application Center. The rate were divided into 4 groups as C (Controls), E (Exercise), T (Trenbolone), and TE (trenbolone + Exercise). The trial period lasted 4 weeks. Supply, care, feeding, and experimental applications of rats were performed in the Selcuk University Experimental Medical Research and Application Center. Anterior and posterior extremities’ bones were dissected and exposed, and the humerus and femur bones exposed were dried. Length, corpus thickness, cortex thickness, and medulla diameter points were determined and the necessary measures were taken. The results are expressed as mean ± SD. ANOVA and Duncan tests were used for the comparison of data. p<0.05 values were considered statistically significant. The mean femoral length was found as 31.31 ± 0.69 in the rats in Group T, 31.46±0.72 in Group E, 31.51±0.58 in Group TE, and 31.48 ± 0.71 in Group C (controls). Examining the mean femoral lengths of Groups T, E, TE and C; the mean femoral length in Group T was numerically higher than that of the Groups E, TE and C, although the difference was not statistically significant (F:0.112; p:0.637). The mean humerus length was found as 24.93 ± 0.59 in the rats in Group T, 24.96±0.68 in Group E, 25.33±0.81 in Group TE, and 25.29±0.77 in Group C (controls). Examining the mean humerus lengths of Groups T, E, TE and C; the mean humerus length in Group T was numerically higher than that of the Groups E, TE and C, although the difference was not statistically significant (F:0.608; p:0.355). We found that the mean values of corpus and cortex thickness, and medullary diameter points were similar in the Groups T, E, TE, and C, and the differences were not statistically significant (p>0.05). Results of this study indicate that trenbolone supplement may lead to early epiphyseal closure in femur and humerus bones of rats, ceasing the increase in their length. We believe that the results obtained from this trenbolone trial will provide important data to the studies that will be conducted on anabolic androgenic steroids.

Key words: Anabolic Androgenic Steroid, Trenbolone, Rat, Femur, Humerus

INTRODUCTION

In the most universal definition, doping is defined as the use, consume or illegal intake of substances prohibited under the rules in order to increase the athletic (12).

Anabolic androgenic steroids (AASs) are mostly used for their anabolic effect, and for increasing endurance and sportive performance by increasing muscle mass and muscular tissue (2, 6).

Frequently preferred anabolic androgenic steroid by athletes and other persons is trenbolone (20). Trenbolone hormone was produced for the first time at the end of 1960s. 19-nortestosterone (19-nor) classification shows structural variance from testosterone since it is deprived of one carbon atom. This difference provides trenbolone to be in the same category with Deca Durabolin (Nandrolone Decanoate) (14). Trenbolone is a highly potent anabolic steroid and primarily preferred by many athletes (20).

Similar to the other steroids, trenbolone significantly increases protein synthesis and nitrogen involvement in muscular tissue. Protein synthesis is rate of cell to produce proteins; protein represent primary constituent of a cell (19). Another steroid feature of trenbolone is inhibition of glucocorticoid hormones. Glucocorticoid hormones that are named as stress hormones in some sources function differently from anabolic steroids in many aspects since these hormones destroy muscular tissue and increase adipose tissue (4).
The objective of this study was to determine the effect of trenbolone supplement administered for four weeks in extremity bones of rats.

**MATERIAL & METHODS**

This study was conducted on 28 rats (Male, Wistar) of 28 days (61.80 g) supplied from the Selcuk University Experimental Medicine Research and Application Center. The rats were divided into four groups as the Control group (C), Exercise group (E), Trenbolone group (T), and Trenbolone + Exercise group (TE). The experiments lasted for four weeks. Supply, care, feeding, and experimental application were conducted in the Selcuk University Experimental Medicine Research and Application Center. The rats were housed in the experimental animals unit, in the plastic rat cages at 23±2°C room temperature and 50±10% relative moist environment, 12/12 light/dark cycle and with feeding ad libitum. Daily freshened water (~ 50 mL/day/rat) was kept available in front of the rats for drinking any time. The study was approved by the Selcuk University Experimental Medicine Research and Application Center Ethical Committee (Decision no: 2018-2, Date: 24/01/2018). The animals were grouped as follows:

- **Group 1**, C (Control) group (n:7): Rats in this group were given standard pellet feed and drinking water ad libitum.
- **Group 2**, E (Exercise) group (n:7): Rats in this group were given standard pellet feed and drinking water ad libitum. The rats were exercised on treadmill at a rate of 25 m/min, 45 minutes a day, 5 days a week for 4 weeks.
- **Group 3**, T (Trenbolone) group (n:7). Rats in this group were given standard pellet feed and drinking water ad libitum during the study. Trenbolone enanthate at a dose of 10 mg/Kg/rat (10) was diluted in 100 mcl peanut oil and administered as intraperitoneal one day a week for four weeks.
- **Group 4**, TE (Trenbolone + Exercise) group (n:7). Rats in this group were given standard pellet feed and drinking water ad libitum during the study. Trenbolone enanthate at a dose of 10 mg/Kg/rat (10) was diluted in 100 mcl peanut oil and administered as intraperitoneal one hour before the exercise, one day a week. The rats in this group were exercised for four week.

Trenbolone Supplement: Trenbolone enanthate (TRENBOLONE E200, Pharma Generics) at a dose of 10 mg/Kg/rat was diluted in 100 mcl peanut oil and administered as intraperitoneal in rats in the groups T (Trenbolone) and TE (Trenbolone + Exercise) for four weeks. Body weight of the rats was measured at the beginning of the study and on the same day during 4 weeks to adjust the weekly dosage (10 mg/Kg/rat) for trenbolone administration.

**Exercise Program**: 8-track treadmill, specially designed for rats was used in exercise application. After an adaptation period of one week (5 days), rats in the exercise groups were exercised on the treadmill at a rate of 25 m/min (1.5 Km/hour) for 45 minutes, 5 days a week over 4 weeks.

**Adaptation protocol:**
- Day 1: 10 m/min, 10 minutes
- Day 2: 20 m/min, 10 minutes
- Day 3: 25 m/min, 10 minutes
- Day 4: 25 m/min, 20 minutes
- Day 5: 25 m/min, 30 minutes

Measurements: At the end of the study, front and back extremities of the subjects were exposed and dissected. Length, corpus thickness, cortex-cortical bone thickness and medullary diameter-cavum medullare measurements were carried out in the exposed humerus and femur bones, using a 0-100 mm caliper.

Anatomic reference points [A (length), B (corpus), C1-C2 (cortex-cortical bone thickness-substantia compacta) and D (medullary diam-cavum medullare)] of the humerus and femur bones at the right side to be measured were determined and the necessary measurements were made in each of these points with a 0-100 mm caliper (Stainless hardened digital caliper, China) (Images 1, 2).

Status of epiphysis was examined in the relevant bones. Images of the bones were taken with a digital camera (Nikon D200, China) (Images 1, 2). In addition, final mean body weight was measured in all subjects with a precision scale before euthanasia.

**Statistical Analysis**: Statistical evaluation of the data was performed utilizing SPSS 18.0 (SPSS 18.0 for Windows/SPSS Inc, Chicago, USA). The results were expressed as mean ± standard deviation. Comparison of the data between the groups was made using ANOVA and Duncan tests. p<0.05 values were considered statistically significant.

Image 1. Reference points of length (A), Corpus (B1+B2/2), Cortex (C1+C2+C3+C4/4) and medullary diameters (D1+D2/2) of humerus of the rats (Right medial side)
A: Distance between the end points of caput humeri and trochlea humeri
B: Corpus thickness of the humerus (lower border level of Tuberositas deltoidea)
C1-C2: Mean femur thickness of the humerus at cortex level (cortical bone-substantia compacta)
D: Cavum medullare diameter of the humerus at cortex level

**Table 1.** Comparison of the mean length, and diameters of corpus, cortex and medullary of femur bones in (Trenbolone), E (Exercise), TE (Trenbolone + Exercise) and C (Control) groups (mm) (mean ± SD).

|        | T     | E     | TE    | C     | Test value, p |
|--------|-------|-------|-------|-------|---------------|
| Length | 31.31±0.69 a | 31.46±0.72 a | 31.51±0.58 a | 31.48±0.71 a | F: 0.112 p:0.637 |
| Corpus | 3.74±0.15 a  | 3.76±0.18 a  | 3.92±0.13 a  | 3.74±0.15 a  | F: 2.797 p:0.062 |
| Cortex | 0.614±0.04 a | 0.605±0.03 a | 0.612±0.04 a | 0.610±0.05 a | F: 0.043 p:0.988 |
| Medullary | 1.987±0.14 a | 2.135±0.20 a | 2.125±0.12 a | 1.975±0.21 a | F: 1.811 p:0.172 |

*Different letters (a,b) at the same row indicate statistical significance (p<0.05)
**F:One way ANOVA/Duncan

The mean femur length of the rats was found as 31.31±0.69 in Group T, 31.46 ± 0.72 in Group E, 31.51 ± 0.58 in Group TE, and 31.48±0.71 in Group C. When the mean femur length values of the Groups T, E, TE and C were examined; the mean length was numerically shorter in Group T compared to Groups E, TE, and C, although the difference was not statistically significant (F:0.112, p:0.637).

The mean femur corpus thickness of the rats was found as 3.74±0.15 in Group T, 3.76±0.18 in Group E, 3.92±0.13 in Group TE and 3.74±0.15 in Group C. When the mean femur corpus thickness values were evaluated; it was found that the mean femur corpus thickness was similar among all groups and no statistically significant difference was found between them (F:2.797, p:0.062).

The mean femur cortex thickness of the rats was found as 0.614±0.04 in Group T, 0.605±0.03 in Group E, 0.612±0.04 in Group TE and 0.610±0.05 in Group C. When the mean femur cortex thickness values were evaluated; it was found that the mean femur cortex thickness was similar among all groups and no statistically significant difference was found between them (F:0.043, p:0.988).

The mean femur medullary diameter of the rats was found as 1.987±0.14 in Group T, 2.135±0.20 in Group E, 2.125±0.12 in Group TE and 1.975±0.21 in Group C. When the mean femur medullary diameter values were evaluated; it was found that the mean femur medullary diameter was similar among all groups and no statistically significant difference was found between them (F:1.811, p:0.172)
Table 2. Comparison of the mean length, and diameters of corpus, cortex and medullary of humerus bones in (Trenbolone), E (Exercise), TE (Trenbolone + Exercise) and C (Control) groups (mm) (mean ± SD).

|       | T       | E       | TE      | C       | Test value, p |
|-------|---------|---------|---------|---------|---------------|
| Length| 24.93±0.59 a | 24.96±0.68 a | 25.33±0.81 a | 25.29±0.77 a | F:0.608 p:0.355 |
| Corpus| 2.47±0.06 a  | 2.51±0.12 a  | 2.52±0.13 a  | 2.46±0.07 a  | F:0.496 p:0.689 |
| Cortex| 0.594±0.06 a | 0.591±0.03 a | 0.594±0.06 a | 0.595±0.02 a | F:0.013 p:0.998 |
| Medullary| 1.51±0.12 a | 1.72±0.14 a | 1.71±0.21 a | 1.52±0.18 a | F:0.246 p:0.652 |

*Different letters (a,b) at the same row indicate statistical significance (p<0.05)
**F:Oneway ANOVA/Duncan

The mean humerus length of the rats was found as 24.93±0.59 in Group T, 24.96 ± 0.68 in Group E, 25.33 ± 0.81 in Group TE, and 25.29±0.77 in Group C. When the mean femur length values of the Groups T, E, TE and C were examined; the mean humerus length was numerically shorter in Group T compared to Groups E, TE, and C, although the difference was not statistically significant (F:0.608, p:0.355).

The mean humerus corpus thickness of the rats was found as 2.47±0.06 in Group T, 2.51±0.12 in Group E, 2.52±0.13 in Group TE and 2.46±0.07 in Group C. When the mean humerus corpus thickness values were evaluated; it was found that the mean humerus corpus thickness was similar among all groups and no statistically significant difference was found between them (F:0.496, p:0.689).

The mean humerus cortex thickness of the rats was found as 0.594±0.06 in Group T, 0.591±0.03 in Group E, 0.594±0.06 in Group TE and 0.595±0.02 in Group C. When the mean humerus cortex thickness values were evaluated; it was found that the mean humerus cortex thickness was similar among all groups and no statistically significant difference was found between them (F:0.013, p:0.998).

The mean humerus medullary diameter of the rats was found as 1.51±0.12 in Group T, 1.72±0.14 in Group E, 1.71±0.21 in Group TE and 1.52±0.18 in Group C. When the mean humerus medullary diameter values were evaluated; it was found that the mean humerus medullary diameter was similar among all groups and no statistically significant difference was found between them (F:0.246, p:0.652).

DISCUSSION

When femur and humerus lengths of Groups T (Trenbolone), E (Exercise), TE (Trenbolone + Exercise) and C (Control) were examined; the mean femur length was numerically shorter in Group T than Groups E, TE, and C, although the difference was not statistically significant (p>0.05).

Studies conducted on experimental animals have reported many side effects of AASs (1, 3, 8, 13, 15, 17, 21). In a study investigating effects of trenbolone application on the urinary system, it was concluded that trenbolone application has a partial effect on the urinary system in experimental group (9).

In a study examining effects of trenbolone administered at different time periods and different doses on female rats, trenbolone was found to cause an increase in density of amniotic fluid in pregnant rats (8).

In a study evaluating effects of trenbolone application on muscles, bones, adipose tissue and hemoglobin levels of rats, it was concluded that trenbolone increased density of muscle and bone tissues (16).

In a study investigating effect of trenbolone on bone mineral density, intramuscular trenbolone application was found to increase bone mineral density without changing hemoglobin density (11).

In another study investigating effects of exercise plus nandrolone supplement on heart muscle of rats, it was reported that heart muscle of the rats administered nandrolone supplement was damaged (7).

In a study in which rats were given AAS for 15 days and effects of AAS in cognitive function of rats were investigated, it was reported that rats in AAS group developed learning and memory disorder (18).

In a study examining effect of trenbolone and testosterone supplement on skeletal muscle growth of rats, it was concluded that testosterone grew skeletal muscle at a higher rate than trenbolone (22).
In a study investigating effects of androgen application on bones of rats, it was reported that femur bones of the rats given androgen remained short (5).

In the present study, when corpus and cortex thickness and medullary diameter values of femur and humerus were examined in T (Trenbolone), E (Exercise), TE (Trenbolone + Exercise) and C (Control) groups; corpus and cortex thickness and medullary diameter values of femur and humerus were similar between Groups T, E, TE and C (p>0.05).

CONCLUSION

Femur length was shorter in the rats in Group T compared to the rats in Groups E, TE and C after trenbolone supplement.

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