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THE RELATIONSHIP BETWEEN POWER OUTPUT, OXYGEN

CONSUMPTION AND HEART RATE DURING EXERCISE IN

THE RAT

DEGREE FOR WHICH THESIS WAS PRESENTED ......
YEAR THIS DEGREE GRANTED .......

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# THE RELATIONSHIP BETWEEN POWER OUTPUT, OXYGEN CONSUMPTION AND HEART RATE DURING EXERCISE IN THE RAT 

by


C PHILIP D. CAMPAGNA III

A THESIS

SUBMITTED TO THE FACULTY OF GRADUATE STUDIES AND RESEARCH IN PARTIAL FULFILMENT OF THE REQUIREMENTS FOR THE DEGREE OF DOCTOR OF PHILOSOPHY


## UNIVERSITY OF ALBERTA

## FACULTY OF GRADUATE STUDIES AND RESEARCH

The undersigned certify that they have read, and recommend to the Faculty of Graduate Studies and Research, for acceptance, a thesis entitled THE RELATIONSHIP BETWEEN POWER OUTPUT, OXYGEN CONSUMPTION AND HEART RATE DURING EXERCISE IN THE RAT submitted by PHILIP D. CAMPAGNA
in partial fulfilment of the requirements for the degree of Doctor of Philosophy in Physical Education.

## DEDICATION

To my wife Marilyn and late Mother Francis
for their moral support over the years
(1)

The power output achieved in and metabolic adjustments $\left(\mathrm{VO}_{2}\right)$ to a wide variety of exercise intensities in rats of different aerobic fitness levels was investigated. Thirty male Wistar rats were divided into five groups; sedentary control (SC), exercisedrun wheel (El), exercised (E2), conditioned-run wheel (Cl) and conditioned (C2). The animals were trained and/or conditioned on a vertical treadmill for 8 weeks. The SC group was trained for a period of l-2 mins., once a week, while the El and E2 groups were exercised twice a week with a progressive increase in duration. The Cl and C2 groups were conditioned four days a week to a maximum of 30 minutes per day. An intermittent test for maximal oxygen consumption was performed at the end of the 7 th and 8 th week. Expired gases were collected from a nose cone and heart rates recorded. There was no significant difference in maximal oxygen consumption between groups. However, the slope of the linear regression lines for $\mathrm{VO}_{2}$ and power output was steeper ( $\mathrm{p}<0.05$ ) for the conditioned (Cl and C2 pooled) as compared to the non-conditioned (El, E2 and SC pooled) groups. This suggests that an adaptation in the conditioned group has occurred. At the lower power outputs, the rats' heart rate increased rapidly to near maximal values. Since $\mathrm{VO}_{2}$ continued to rise at power output levels above the levels where heart rate has levelled off, the increased $\mathrm{VO}_{2}$ must be achieved by enhanced extraction at the tissue level. Therefore, the different slopes in conjunction with similar maximum $\dot{V}_{2}$ values for the different groups suggest that the rat may not be an ideal model for man when studying exercise adaptations in the oxygen transport system.

To Dr. Howard A. Wenger, Ph.D., I express my gratitude for his advice and help over the past few years.

To Dr. James C. Russell, Ph.D., a very special thank you for the many hours of guidance and support, I express my deepest appreciation.

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For over 100 years, the domesticated rat has been used for scientific studies in Europe. The Norway rat (Rattus norvegicus) was the first animal to be domesticated for the sole purpose of scientific research (Richter, 1968). Adolf Meyer imported the domesticated rat to North America for studies on the brain in 1890 and shortly thereafter, H.H. Donaldson recognized their potential for scientific research and established the famous Wistar colony in Philadelphia (Richter, 1968).

The use of animals such as the laboratory rat in research allows a range of manipulation of both dependent and independent variables, which is often inconvenient or impossible, either for practical or ethical reasons, when studying humans (Plaut, 1975). Experimental variables of interest can be manipulated a great deal with animal populations along with some other factors which otherwise might affect the outcome of the study. Ready access to tissue samples e.g. brain, heart, liver, etc. from the rat are not normally permitted when researching with humans (Plaut, 1975; Lane-Petter, 1976). Furthermore, the undertaking of longitudinal studies as well as replication and the extension of previous work is made possible because of the short-life span (approx. 3 years) and growth period of the rat (Richter, 1968; Plaut, 1975; Roberts and Goldberg, 1976). Of particular advantage the rat's size and economy as well as its high resistance to infection, makes it an ideal model for surgical and physiological studies (Richter, 1968; Plaut, 1975).

The laboratory rat has been used for some time in the
medical sciences. It's dietary needs are very similar to those of man, thereby providing discoveries of the nutritive value of vitamins (Matschiner et al., 1967; Bieri et al., 1969). Various strains of laboratory rats have also been bred in order to study cardiovascular diseases (Okamoto, 1964), hypertension (Koletsky, 1973) and obesity (Bray, 1977). The rat has many other medical applications such as clinical investigation of diseases, cancer research, drug research as well as the study of endocrine function and environmental factors.

Until recently, research in exercise physiology has been conducted solely on humans. However, this situation has changed considerably over the past ten years with animal research now being carried out frequently (Oscai and Mole, 1975). A substantial number of studies have examined the acute and chronic effects of conditioning on enzyme activities (Saubert et al.,1973; Hollozy et al., 1975; Hickson, 1976; Jobin, 1976), ligaments and tendons (Tipton et al., 1975; Booth and Gould, 1975), muscular hypertrophy (Goldberg et al., 1975; Baldwin et al., 1977), temperature regulation (Baker and Horvath, 1964; Gollnick and Ianuzzo, 1968; Popovic et al., 1969), hormones (Trenkle, 1974; Gollnick and Ianuzzo, 1975), proteins and nucleic acid concentrations (Wilkinson, 1977), muscle structure (Bowers, 1974; Cosmas and Edington, 1975) and many other variables using the rat as a model.

An underlying assumption when dealing with animal models such as the rat is that it is possible to obtain data on animals
and then extrapolate these findings to humans (Oscai and Mole, 1975). In many cases however, generalizability from animal studies may not always be possible and therefore may just indicate research needed with human subjects (Hardin, 1965; Oscai and Mole, 1975). If exercise physiologists are to extrapolate to man the data which has been obtained on the laboratory rat during exercise, then it is necessary to show similarities between the rat and man. Only in this way can the rat be thereby justified as an adequate model for research in this area.

Although the resting heart rate of the rat is approximately five times that of man, research has shown that bradycardia with endurance conditioning occurs in the rat (Barnard et al., 1976; Tipton, 1965; Lin and Horvath, 1972) as well as in man (Sutton et al., 1967; Frick et al., 1967; Ekblom et al., 1973). The heart rate (Jones et al., 1960) and metabolic adjustment (Shepherd and Gollnick, 1976) of the rat to submaximal steady state exercise is very similar to the pattern observed in man (Mathews and Fox, 1976). Shepherd and Gollnick (1976) have shown a linear increase in oxygen consumption with work intensity (running speed) up to maximum in the rat. This relationship is well established in humans (Astrand and Rodahl, 1970; Mathews and Fox, 1976). Barnard et al. (1974) reported a decrease in maximum heart rate with aging in the rat, which corresponds to the decrease in maximum heart rate seen in man with increasing age (Shepherd, 1968; Robinson et al., 1976).
the rat were significantly higher than the rates recorded at a submaximal endurance run (Barnard et al., 1974). This is in agreement with Saltin's (1964) data for humans. A graded treadmill test for rats developed by Wranne and Woodson (1973) showed a linear increase in heart rate with an increase in running speed and/or gradient; a well known phenomenon in man (Astrand and Rodahl, 1970; Mathews and Fox, 1976). These results are contrary to the findings of Barnard et al. (1974) who observed heart rate responses to various levels of exercise which are quite different than the response observed in humans. A closer look at their data (Barnard et al., 1974) revealed some interesting findings. In calculating the percentage of maximum heart rate at submaximal levels, they suggested that the relative change in heart rate is from zero beats per minute to the submaximal heart rate and the range of maximum heart rate is from zero to maximum. In doing so, they are implying that the range between zero and resting is a functional portion of the increase and therefore inflate the calculated percentages. Wranne and Woodson (1973) on the other hand used the relative change from resting heart rate to submaximal heart rate and resting heart rate to maximum heart rate as their functional ranges. Therefore, their (Barnard et al., 1974) reported value of 80\% maximum heart rate at the lowest power output (13.4 meters/min.) is dramatically reduced to $50 \%$ when utilizing the functional range of Wranne and Woodson. Thus, with application of Wranne and Woodson's functional range to the data of Barnard et al. (1974) a linear relationship between heart rate and running speed exists.

Blood lactate levels at maximum work for the rat (Wranne and Woodson, 1973) resemble concentrations that are observed in man at maximal work (Astrand and Rodahl, 1970). According to Popovic et al. (1969) the stroke volume of the rat during exercise does not increase. Similar results are reported in man when muscular work is performed in the supine position (Edington and Edgerton, 1976), the heart is in a similar position in the rat during exercise on the horizontal treadmill.

Although the rat's capacity to dissipate heat appears to be poor, a graded effect on body temperature during exercise to exhaustion (Gollnick and Ianuzzo, 1968) corresponds to observed values reported for humans (Robinson, 1963).

Along with the changes that occur in the cardiovascular system, major biochemical adaptations can occur in skeletal muscle with conditioning. Mammalian skeletal muscles have been classified by a number of different systems during the past fifteen years (Close, 1972). The classification scheme utilized by Peter et al. (1972) seems to be the most functional. They classify muscle fibers as fast-twitch-glycolytic (FG), fast-twitch-oxidative-glycolytic (FOG) and slow-twitch-oxidative (SO) according to contractile speed and metabolic properties. There are a few basic metabolic differences among the skeletal muscles of the rat and human. In man, the SO muscle fiber has the highest oxidative potential, while the FOG has an intermediate oxidative potential lying somewhere between the SO and FG fibers (Holloszy et al., 1975). The SO muscle fiber of the rat on the other hand has intermediate
oxidative potential with the FOG fiber being the most oxidative (Holloszy et al., 1975). It should be pointed out that the fiber composition of human skeletal muscle is heterogeneous in nature. Whereas, in the rat a number of muscles such as the soleus and plantaris are relatively homogeneous in their fiber composition. Although this appears to be a disadvantage in attempting to extrapolate from rat to man, it allows the researcher an opportunity to examine the relative effect of various conditioning protocols on certain fiber type populations (Holloszy and Booth, 1976).

Holloszy (1967; 1973) has shown that major biochemical adaptations occur in skeletal muscle of the rat with endurance conditioning. A finding of similar biochemical adaptations has been confirmed by Morgan et al. (1971) for human subjects. Depletion of glycogen stores has been reported to occur in both rat (Gollnick et al., 1970) and man (Lappage, 1969; Taylor, 1971) during prolonged exercise. Free fatty acid utilization and uptake as well as their release from adipose tissue has also been shown to be influenced by prolonged activity in both rat (Jones and Havel, 1967) and man (Hagenfeldt and Wahren, 1971). Many investigators have reported an increase in skeletal muscle glycogen content for rats (Grollman, 1955; Gollnick et al., 1970) and man (Bergstrom and Hultman, 1966; Lappage, 1969) following regular conditionịng programs. An increase in total mitochondrial protein due to increases in the size and number of mitochondria, has been reported in skeletal muscle in both man and rats (Holloszy and Booth, 1976).
presented, it is suggested that the laboratory rat is an excellent experimental model for research in exercise physiology.

Exercise physiologists have employed running and swimming programs most frequently to evoke a conditioning response. The motordriven treadmill (Critz and Merrick, 1962; Andrews, 1965; Bhatia et al., 1966; Koniski, 1966, Brannon et al., 1968), running wheel (Wells and Heusner, 1971) and exercise drum (Jenner and Byrd, 1974; Shepherd and Gollnick, 1976) are the most common modes for running the laboratory rat. Many laboratories use swimming as a form of exercise with the addition of weights as a means of standardizing the workload (Dawson and Horvath, 1970).

Quantification of the work done by each animal while swimming is unknown and may be influenced by a number of factors (Griffiths, 1960; Baker and Horvath, 1964a; McArdle, 1967; Hardin, 1968). Treadmill running, exercise drums and running wheels provide some control over the speed at which the animals are to train, however, because of the horizontal component the amount of work or power output actually performed is difficult to calculate in physical terms (Hardin, 1965).

Researchers have employed various conditioning protocols over the past ten years and have shown a number of physiological changes. Table $l$ represents a number of conditioning protocols that have been reported in the literature. A few authors (Baldwin et al., 1977) have made an attempt to justify the conditioning protocol used, however, the majority give no rationale for the selection of the speed, incline, frequency and duration. Often
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TABLE 1 A SELECTION OF CONDITIONING PROGRAMS UTILIZED BY RESEARCHERS

| $\begin{gathered} \text { AUTHOR } \\ \& \\ \text { YEAR } \end{gathered}$ | SPEED $(\mathrm{m} / \mathrm{min})$ | INCLINE <br> (deg) | FREQUENCY (days/wk) | WORK TIME (mins/day) | DURATION (weeks) | APPARATUS |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Bagby, 1972 | 26.8 | ? | 5 | 60 | 11 | treadmill |
| + Edington, 1973 | 30 | 8 | 5 | 60 | 16 | treadmill |
| Taylor, 1974 | 26.8 | ? | 5 | 60 | 12 | treadmill |
| \# Borensztajn, 1975 | 31 | 8 | 5 | 120 | 12 | treadmill |
| Barnard, 1976 | ? | ? | 5 | 60 | 12 | treadmill |
| Hickson, 1976b | 36 | - | 5 | 50 | 8 | run-wheel |
| Terjung, 1976 | 26.8 | 5.7-8.5 | 5 | 60 | 12-18 | treadmill |
| Wachtova, 1976 | 18-20 | ? | ? | 120 | 7.6 | treadmill |
| * Baldwin, 1977 | 26.8 | 14 | 5 | 60 | 8 | treadmill |
| Fitts, 1977 | 32.2 | 8.5 | 5 | 120 | 27 | treadmill |
| Pitts, 1977 | 18 | 8 | 7 | 60 | 15 | treadmill |

[^0](2)
important information is not reported, thereby making comparisons between studies extremely difficult (see Table l).

If the primary objective of a conditioning program is to achieve optimal improvements in a selected variable then appliction of the progressive overload principle is essential (Mathews and Fox, 1976). With endurance conditioning the intensity of the program rather than the duration has been reported to elicit the greatest improvements (Shepherd, 1969; Wenger and Macnab, 1975). Astrand and Rodahl (1977) have stated that "an adaptation takes place to a given load; in order to achieve further improvements, the training intensity has to be increased."

A common procedure is to progressively increase the speed of the treadmill and duration of the run during the first few weeks and then to maintain the animals at a selected speed and duration throughout the remainder of the experiment. Whether the procedures employed by most researchers (see Table l) progressively overload the rat throughout the entire duration of the study is questionable. If the intent is to compare the adaptation of conditioning and/or exercise between the rats of various studies or to extrapolate to man, then the relative intensity of the conditioning program should be known. The determination of the precise power output the rat is performing on the horizontal treadmill is seriously confounded by the horizontal component of running and/or gait changes.

A vertical treadmill has been designed which allows the power output performed by the rat to be measured (Russell et al.,
1978). Along with the measurement of oxygen consumption (considered the best measure of aerobic metabolic rate) researchers will be able to determine the relative intensity of their conditioning programs.

Thus, if mechanical and metabolic work as well as the heart rate responses to different loads can be determined, it will enhance the capability to quantify the conditioning stimulus. This will in turn improve the generalizability to humans during exercise.

The purpose of the study is:
To determine the relationship between the power outputs achieved in and heart rate and metabolic adjustments $\left(\dot{\mathrm{VO}}_{2}\right)$ to a wide variety of conditioning intensities in rats of different aerobic fitness levels.

Terminology

Maximal oxygen consumption - the maximal rate at which oxygen can be consumed/minute.

Power output - performance of work expressed per unit of time. Training - activity designed to teach a particular skill. Exercise - any physical activity not intended to improve skill, strength or endurance, but to maintain these.

Conditioning - activity designed to improve the physical performance such as strength or endurance, as opposed to skill.

ANIMAL SELECTION AND CARE

Thirty male Wistar rats ( WOF (WI) Specific pathogen free CFN) approximately five to seven weeks of age were obtained through the Director of the Health Sciences Animal Center at the University of Alberta from the Woodlyn Farms, Guelph, Ontario. Upon arrival, the animals were randomly assigned to one of five groups; a control group (SC), two exercised groups (El and E2) and two conditioned groups (Cl and C2). The control group (SC) was essentially given no exercise, although they were exposed to the treadmill once a week. The exercised group El was exercised on the treadmill twice a week and had voluntary access to a running wheel (see Fig. 1). Whereas,


FIGURE 1 VOLUNTARY RUNNING WHEEL

TABLE 2

## DESCRIPTION OF EXPERIMENTAL GROUPS AND NUMBER OF ANIMALS ASSIGNED TO EACH CONDITION

ANIMAL GROUP
NUMBER OF ANIMALS
(N)

```
SC}\mp@subsup{}{}{1}\mathrm{ - Sedentary control group
1 0
    (no exercise)
El - Exercised group (exercised on treadmill
5
    twice a week, access to running wheel)
E2 - Exercised group (exercised twice a week
    5
        on a treadmill)
Cl - Conditioned group (conditioned 4 d/wk.
        on a treadmill, access to running wheel)
C2 - Conditioned group (conditioned 4 d/wk.
5
        on a treadmill)
```

$1_{\text {Although }}$ this group is designated as receiving no exercise, they were exposed to the treadmill once a week.
exercised group E2 was exercised only twice a week on the treadmill.
Both conditioned groups $C l$ and $C 2$ were conditioned on the treadmill
four days a week, however the Cl group had voluntary access to a
running wheel (see Table 2). The animals were housed in standard
$9^{\prime \prime} \mathrm{x} 18^{\prime \prime} \mathrm{x} 6^{\prime \prime}$ cages. Animals in groups El and Cl (Table 2) were exposed to the running wheels on alternate days because of a limited number of running wheels.

Purina rat chow containing $23 \%$ crude protein and water was provided ad libitum. All animals were given a week to adapt to the new environment which included a reversal of the day-night cycle (light 6 pm . to $6 \mathrm{am}$. ). Weighing of all animals in groups SC, E2
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and C2 was carried out every 3 to 5 days (see Appendix B-1). Animals in groups El and Cl were weighed every day (see Appendix B-l). Each morning, the cages were rotated on the cage rack, soiled papers in the running wheels were changed and food and water replenished.

## DESCRIPTION OF TREADMILI

The animals were trained and conditioned on a vertical treadmill (rotating ladder) consisting of $3 / 8^{\prime \prime}$ roller chains, l.ll meters in length, connected by $3 / 10^{\prime \prime}$ rods with a $3 / 4^{\prime \prime}$ pitch (see Fig. 2 and Fig. 3). The ergometer is powered by a General Electric D.C. variable speed motor which allows the experimeter a constant speed range of 5.7 to 17.4 meters per minute. Small variations in vertical elevation are possible with a range of $69^{\circ}$ to $90^{\circ}$ (see Appendix C). Motivation to run was provided by a compressed air jet, which was hand-held at the bottom of the treadmill.

TRAINING PROTOCOL

At the end of the adaptation week, all animals were given seven days of orientation to climbing on the vertical treadmill. Following this brief orientation, the maximum heart rate was determined for each animal by progressively increasing the speed until no further increase in heart rate occurred. For the remainder of the study, the sedentary control group (SC) was trained for a period of $1-2$ minutes once a week (Sat.), while the exercise groups El and E2 were exercised twice a week (wed. and Sat.) with a progressive increase in duration (see Appendix D-1). The conditioned groups Cl and C2 were conditioned four days a week (Mon., Tues., Thurs. and
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Figure 2 Front view of vertical treadmill

Figure 3 Vertical treadmill and equipment for the measurement of expired gases (A, vertical treadmill; $B$, Elow meter; $C$, mixing chamber; $D$, vacuum pump)


Fri.).

Each conditioning day consisted of two sessions of progressive duration up to a maximum of fifteen minutes for most animals (see Appendix D-2). The morning sessions were held between 0700 and 1000 hours and the afternoon sessions between 1400 to 1700 hours.

In an attempt to progressively overload the animals in groups Cl, C2, El and E2. a criterion heart rate from 580 to 610 beats per minute was chosen for each animal. As the maximum heart rate decreased with aging, the criterion heart rate was reduced to a range of 570 to 590 beats per minute. The speed of the treadmill was therefore adjusted to maintain the animal at the criterion heart rate (see Appendix H) so that the same relative intensity was maintained throughtout.

Since precise measurements of power output will enhance the capability to quantify the training stimulus, a pilot study was carried out in an attempt to determine the steepest incline, animals could manage and still have a sufficient duration of exercise. From $69^{\circ}$ to $76^{\circ}$ there was little difference in running times. However, longer running times were found to occur at $76^{\circ}$ than at steeper grades. The small angular difference allows the rat to exercise on the treadmill without having his center of gravity outside his hind feet. Therefore, the overhead work of maintaining this position is minimized. The vertical distance travelled is determined by finding the sine of angle theta $(\theta)$ times the distance travelled (Mathews and Fox, 1976). The sine of $76^{\circ}$ is 0.97 , however

increasing the incline to $90^{\circ}$ only increases the sine $\theta$ to 1.00 . Therefore from a physical point of view, $76^{\circ}$ will allow precise measurements of vertical distance travelled as well as provide a sufficient duration of exercise.

EXPERIMENTAL DESIGN

In the present study, five groups of experimental animals were selected (Table 2). The number of training and conditioning days, access to running wheels (for some animals) and a variety of conditioning intensities were employed in an attempt to obtain a wide range of maximal oxygen consumption values.

An intermittent test for maximal oxygen consumption was performed at the end of the seventh week. The duration of the study depended upon the animals reaching asymptote of maximal oxygen consumption, inspite of additional conditioning. A levelling off criterion of $5 \%$ or less over a one week period was established for termination of the study. At the end of the eighth week, all but two animals showed a levelling off of maximal oxygen consumption. The remaining 2 animals showed a levelling off at the end of the nineth week.

At the end of the nineth week, three animals were trained to run on a horizontal treadmill (Collins) consisting of a wide endless belt on rollers divided into 5 compartments. Motivation to run was provided by a compressed air jet, winich was hand-held at the back of the compartment. Arter three consecutive training days, a submaximal oxygen consumption test was performed. The animals ran at a variety of speeds and inclinatıons for a duration of three minutes

each (Table 4).

DETERMINATION OF HEART RATE

During the week of adaptation, two electrodes were placed on each animal for determination of heart rates. The rats were anesthetized lightly with halothane using a Fluotec 3 vaporizer. Two stainless steel 3-0 monofilament sutures were placed bilaterally on the back of the animal approximately one inch from the dorsal midline and just above the scapula (Eisentein and Woskow, 1958).

When heart rates were recorded, two Mueller \#34c alligator clips soldered to a length of Belden \#8429-10 twisted cable were clipped to the sutures on the animal's back. EKG paste was applied to the ends of the alligator clips to provide a better conductive medium between the sutures and clips. The leads were then connected to a Sanborn 350-3200A ECG Preamplifier. For the purposes of the conditioning sessions, a storage oscilloscope was connected to the output of the ECG Preamplifier (see Fig. 4).


FIGURE 4 A SCHEMATIC REPRESENTATION OF THE APPARATUS FOR DETERMINATION OF HEART RATE
(1)

Heart rates were recorded during basal conditions, resting in the cage, pre-exercise ana during exercise. Basal heart rates were recorded during the light portion of the daily cycle. If at all possible, care was taken to not awaken the animal during the measurement of basal heart rates.

DETERMINATION OF OXYGEN CONSUMPTION

For the collection of the expired gases, a nose cone (see Fig. 5 and 6) was developed (open system). Its operation depended upon a negative pressure with the air drawn through by a vacuum pump. Air flow was measured by a Matheson 0.200 - 20 liter flowmeter (see Fig. 3). The rate of air flow was set at 5.5 liters per minute. The air samples were immediately analyzed by a Beckman metabolic cart. A 500 ml . aliquot of air was sampled from the mixing chamber (see Fig. 3) and analyzed for $\% \mathrm{O}_{2}$ and $\% \mathrm{CO}_{2}$ by means of an OM-ll oxygen analyzer and a LB-2 carbon dioxide analyzer. Reference gases of a known concentration (Consumer' Welding Supplies Ltd.) were used for calibration of the gas analyzers. The total response time of the system was eight seconds. Calculation of oxygen consumption was corrected for STPD.

Maximal oxygen consumption was determined by a discontinuous treadmill test. The animal climbed for four minutes, followed by a rest period of approximately $10-15$ minutes. The treadmill speed was then increased and the procedure repeated. Expired gases were measured throughout the test until an asymptote occurred or a decrease in oxygen consumption was produced with increased power output.


Figure 5 Collection of pre-exercise oxygen consumptions from a rat on the vertical treadmill

Figure 6 Mask utilized for collection of expired gases


DETERMINATION OF POWER OUTPUT

The power output performed by the animal is reported in SI units (watts). The following equation was applied to calculate the power output (Russell et al., 1978):

$$
P=W \times V \times 9.80 \times 10^{-3} \times \sin \theta
$$

where: $W$ is the animals weight in grams.
$V$ is the ladder velocity in meters per second.
$\theta$ is the angle between the ladder surface and horizontal.

STATISTICAL ANALYSIS

A one-way analysis of variance (Winer, 1977) was carried out on the data to compare groups. The Newman-Keuls test was used as a post hoc procedure to locate significant differences between group data which was pooled into either conditioned or unconditioned groups. Group regression lines were calculated using all data points from each animal (up to $V_{2} \max$ ). A test for homogenity of regression was used to test the slopes of the regression lines for the conditioned (Cl and C2 pooled) and unconditioned (El, E2 and SC pooled) groups. An alternative method for calculating the group regression lines was used (post hoc), where the slopes of the individual regression lines (Appendix E) for each group were averaged (see Addendum). Significant differences were accepted at the alpha level p is less than 0.05 , where $p$ is the probability that no difference exists.
(

The results are summarized in tabular and graphical form representing the mean values and standard error of the mean (SEM) for each group. Raw data of the dependent variables for all experimental animals can be found in Appendix B. Individual regression lines for oxygen consumption ( $1 / \mathrm{min}$ ) and power output for each animal is located in Appendix E. Tables of statistical analysis and post hoc procedures are located in Appendix F.

Body weights increased with growth throughout the experiment with no difference between groups at the beginning of the conditioning program (Fig. 7). The sedentary control (SC) and exercised (E2) group showed the highest body weight at the end of the experiment and were not different from each other ( $\mathrm{p}>0.05$ ). In descending oruer the exercised-run wheel (El), conditioned (C2) and conditioned-run wheel (C2) were different from the former two groups and from each other (p<0.05) after 58 days (Fig. 7).

Maximal oxygen consumption/unit body weight is illustrated in Fig. 8. The exercised group (E2) has a significantly lower maximal oxygen consumption ( $\mathrm{ml} / \mathrm{kg} / \mathrm{min}$ ) than the sedentary control (SC) and the conditioned groups (Cl and C2). There was no difference ( $p>0.05$ ) among groups in maximal oxygen consumption when expressed in $1 / m i n$.

The linear regression lines for the oxygen consumption ( $1 / \mathrm{min}$ ) and power output for each experimental group is shown in Fig. 9. The slope for the conditioned groups (C1 and C2 pooled) was steeper ( $p<0.05$ ) than the slope for the non-conditioned group (SC, El and E2 pooled).
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Table 3 Heart rates at different levels of activity obtained from the rats during the 2 nd and 8 th week of the experiment

|  | $\begin{gathered} \text { Basal } \\ \text { (asleep) } \end{gathered}$ | Resting <br> in cage | Pre-exercise | Maximum Exercise | Paroxysmal <br> Tachycardia |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 2nd week | $\begin{aligned} & 407(6) \\ & \pm 10 \end{aligned}$ | $\begin{aligned} & 447(6) \\ & \pm 21 \end{aligned}$ | $\begin{aligned} & 504(25) \\ & \pm 4 \end{aligned}$ | $\begin{aligned} & 647(25) \\ & \pm 3 \end{aligned}$ | 844 (1) |
| 8th week | $\begin{aligned} & 300(5) \\ & \pm 8 \end{aligned}$ | $\begin{aligned} & 354(4) \\ & \pm 12 \end{aligned}$ | $\begin{aligned} & 403(20) \\ & \pm 5 \end{aligned}$ | $\begin{aligned} & 597(20) \\ & \pm 3 \end{aligned}$ | - |

$\overline{\mathrm{x}} \pm \mathrm{SE}(\mathrm{N})$
There was no significant difference in pre-exercise oxygen consumption ( $\mathrm{ml} / \mathrm{kg} / \mathrm{min}$ ) for the experimental groups (Appendix B).

Heart rate data obtained from rats during the 2 nd and 8 th week of the experiment is shown in Table 3. Maximum heart rate decreased ( $\mathrm{p}<0.01$ ) from $647 \pm 3$ beats $/ \mathrm{min}$. during the 2 nd week to $596 \pm 3$ beats/min. in the 8 th week of the experiment. There was no significant difference in maximal heart rate between the experimental groups at either the 2 nd or 8 th week. Pre-exercise heart rates decreased (p<0.01) from $504 \pm 4$ beats/min. during the 2 nd week to $403 \pm 5$ beats/min. in the 8 th week. There was no significant difference in the pre-exercise heart rate between the experimental groups in either the 2 nd or 8 th week of the experiment.

The resting heart rates recorded in the cages after onehalf hour rest decreased ( $p<0.01$ ) from $447 \pm 21$ beats $/ \mathrm{min}$. in the 2nd week to $354 \pm 12$ beats/min. in the 8 th week. Basal heart rates while the animals were asleep also showed a decrease ( $p<0.01$ ) from $407 \pm 10$ beats $/ \mathrm{min}$. in the 2 nd week to $300 \pm 8$ beats $/ \mathrm{min}$. in the 8 th

week of the experiment.
Figure 10 shows the relationship between heart rate and power output for the different experimental groups. At the lower power outputs, the rats' heart rate increased rapidly to near maximum values and then plateaued.

Horizontal Treadmill Exercise

Submaximal oxygen consumption and heart rate values obtained
while selected animals were running on the horizontal treadmill are shown in Table 4. Oxygen consumption and heart rate increased progressively when speed was held constant and grade elevated or vice versa. The power output was calculated from the vertical component only and does not include the horizontal component of running.

Table 4 Submaximal measurements of three rats at varying running speeds and inclinations on the

| Animal | Weight (grams) | $\begin{aligned} & \text { Speed } \\ & (\mathrm{m} / \mathrm{min}) \end{aligned}$ | Grade (degrees) | Power Output (watts) | Heart Rate (b/min) | $\stackrel{V O}{2}_{(1 / \mathrm{min})}$ | $\begin{gathered} \stackrel{\mathrm{VO}}{(\mathrm{ml} / \mathrm{kg}}\}_{\mathrm{min}} \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| BL3 | 246 | 26.8 | 4 | 0.075 | 529 | 0.0131 | 53.2 |
| PRR | 349 | 24.8 | 4 | 0.106 | 546 | 0.0197 | 56.5 |
| GRI | 343 | 26.8 | 4 | 0.105 | 563 | 0.0156 | 45.5 |
| BL3 | 246 | 26.8 | 10 | 0.187 | 546 | 0.0169 | 68.7 |
| PRR | 349 | 26.8 | 10 | 0.265 | 581 | 0.0201 | 57.7 |
| GRI | 343 | 26.8 | 10 | 0.260 | 584 | 0.0178 | 52.0 |
| BL3 | 246 | 26.8 | 16 | 0.297 | 554 | 0.0192 | 78.1 |
| PRR | 349 | 26.8 | 16 | 0.421 | 590 | 0.0206 | 59.1 |
| GRI | 343 | 26.8 | 16 | 0.413 | 595 | 0.0187 | 54.6 |
| BL 3 | 266 | 30.0 | 4 | 0.090 | 546 | 0.0101 | 38.5 |
| PRR | 357 | 30.0 | 4 | 0.122 | 552 | 0.0204 | 57.2 |
| BL3 | 266 | 30.0 | 10 | 0.224 | 581 | 0.0181 | 68.8 |
| PRR | 357 | 30.0 | 10 | 0.304 | 577 | 0.0250 | 70.0 |
| BL3 | 266 | 30.0 | 16 | 0.355 | 590 | 0.0204 | 77.8 |
| PRR | 357 | 30.0 | 16 | 0.482 | 590 | 0.0278 | 77.7 |
| BL3 | 263 | 35.0 | 4 | 0.104 | 563 | 0.0133 | 51.1 |
| BL3 | 263 | 35.0 | 10 | 0.258 | 581 | 0.0175 | 67.1 |





- $\overbrace{0}^{8}$
- $0^{2}$

0

(1) SC - SEDENTARY CONTROL

## DE2-EXERCISED

## * EL - EXERCISED - RUN WHEEL

## ocr -CONDItioned



| .4 | .5 | .6 | .7 | .8 | .9 | 1.0 |
| :---: | ---: | ---: | ---: | ---: | ---: | :--- |
| POWER OUTPUT | (watts) |  |  |  |  |  |

[^1]

Eody weight progressively increased throughout the duration of the study in all experimental groups (Fig. 7). This observation has been well documented in a variety of conditioning and exercise programs such as isometric conditioning (Zika et al., 1975; Exner et al., 1973a and b), sprint conditioning (Staudte et al., 1973; Houston and Green, 1975; Hickson et al., 1976a), endurance conditioning (Baldwin et al., 1975; Huston et al., 1975; Muller, 1975). Houston and Green (1975) suggested that an appetite-suppressing effect occurs when rats are subjected to a vigorous conditioning program. This effect may combine with an increased energy expenditure thereby resulting in a smaller increase in the body weights of the exercised and/or conditioned groups.

At the conclusion of the present study, the group that exercised once a week (E2) did not differ in body weight from the sedentary control group. However, the other experimental groups had significantly lower body weights than the E2 and SC groups and were also significantly different from each other (Fig. 7). The experimental group exposed to the greatest amount of total work ( C 2 : conditioned-run wheel) displayed the least weight gain followed by the conditioned (Cl) and exercised-run wheel (El) groups respectively. It appears that body weight changes are not only dependent upon the age and sex of the animal but also upon the intensity, duration and total work of the exercise and/or conditioning program.

The most popularly used procedure to evaluate the intensity
or quantity of metabolic overload in an exercise bout has been the measurement of oxygen consumption (Scheur and Tipton, 1977). It reflects the efficiency of uptake, transport and utilization of oxygen by the cardiorespiratory and muscular systems. The maximal oxygen consumption values observed in the present study ranged from 64.4 to $90.7 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. (Fig. 8). The experimental group that exercised (E2) twice a week had a significantly lower maximal oxygen consumption than the sedentary control (SC) and conditioned groups (Cl and C2). There seems to be no apparent reason for the lower values obtained from the E2 group as compared to the sedentary controls. There was no significant difference in maximal oxygen consumption between the other experimental groups in spite of substantially different amounts of chronic exercise. The values reported in this study are slightly lower than those previously reported in the literature for rats running on a treadmill or running wheel. Wilson et al. (1978) reported maximal oxygen consumption values of $91 \pm 2$ and $87 \pm 2 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. for unconditioned male and female rats respectively. Shepherd and Gollnick (1976) reported a maximum value of $95 \pm 1.4 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. for male rats conditioned five days a week for a period of six weeks. A number of possible reasons exist for the higher values reported in the literature. For example, Wistar rats were used in the present study, whereas Shepherd and Gollnick (1976) selected rats from the Sprague-Dawley strain. Wilson et al. (1978) chose Okomato-Aoki as well as Sprague-Dawley rats for their study. The type of ergometer utilized to obtain maximal values was different in each study. Wilson et al. (1978) employed a horizontal treadmill
while Shepherd and Gollnick (1976) used a motor-driven running wheel. In the present study, a vertical treadmill was used which required a climbing motion rather than a running motion. Both Shepherd and Gollnick (1976) and Wilson et al. (1978) failed to report their values corrected to S.T.P.D. (Standard, Temperature, Pressure, Dry, e.g. $760 \mathrm{mmHg} ., 0^{\circ}$ C.). This was likely an oversight. However, if the reported values were not corrected to S.T.P.D., the differences in maximal oxygen consumption between this study and the reported ones would not exist. The maximal oxygen consumption values reported for swimming rats range from 55 to $80 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. (Baker and Horvath, 1964b; Kratzing and Mark. 1967; McArdle, 1967; Dawson and Horvath, 1970). These reported values are lower than the observed values in the present study. It is difficult to compare swimming to running since the physical quantification of the power output while swimming is very difficult. McArdle (1967) has suggested that the addition of weight to the swimming rat does not increase the oxygen consumption but actually reduces the ability of the animal to exchange gases at the surface. The lower values of maximal oxygen consumption obtained while swimming would suggest that swimming lends itself to submaximal exercise programs but is less suitable to overloads of near maximal intensity. Since there was essentially no difference in maximal oxygen consumption between experimental groups, one could conclude that the conditioning program produced no change in the aerobic capacity of the rats. However, when the slopes of the regression lines for the conditioned (Cl and C2 pooled) and non-conditioned (SC, El and E2 pooled) groups are compared, they differ significantly

(Fig. 9). This suggests that an adaptation in the conditioned group has occurred. A possible explanation is that at the higher power outputs the conditioned rats were able to meet the increased demands through increased aerobic catabolism whereas, the nonconditioned animals derived a greater percentage of the required energy to perform the work from anaerobic sources. This could imply that chronic conditioning, as performed in this study, elicits an increased emphasis on aerobic metabolism at higher power outputs rather than an elevated maximal oxygen consumption. In man oxygen consumption at submaximal power outputs are very similar but the conditioned athlete is able to extend his oxygen consumption. The different slopes (also see Addendum) in conjunction with similar maximal oxygen consumption values for the different groups suggests that the rat may not be an ideal model for man when studying exercise adaptations in the oxygen transport system. Pre-exercise oxygen consumptions were higher ( $34 \pm 2 \mathrm{ml} / \mathrm{kg} /$ min.) than most values reported in the literature (Kratzing and Mark, 1967; Popovic et al., 1969; Shepherd and Gollnick, 1976). Since the rats were accustomed to wearing the mask only when placed on the treadmill, it is suggested that they were anticipating the start of exercise. This is consistent with the elevated resting oxygen consumptions reported by Shepherd and Gollnick (1976) and also with the anticipatory rise in colonic temperature of rats prior to exercise reported by Gollnick and Ianuzzo (1968). Also the mask itself may precipitate elevated metabolic rates in contrast to the chambers used traditionally.

The individual regression line for oxygen consumption
and power output indicates that a linear relationship exists (Appendix E). This is consistent with the findings in man (Mathews and Fox, 1976). The adjustment to submaximal steady state exercise in the rat also follows a pattern that is often documented in human research (Astrand and Rodahl, 1977). There was a 3 to 4 minute period before oxygen consumption plateaued and in many cases then showed a drop which is frequently evident in man. Hence, although the response to increasing power outputs is quite different in the rat model, the lag time associated with the slow accomodation in the cardiovascular system to a single power output is comparable to man.

The maximal heart rates observed during the 2 nd and 8 th experimental week (Table 3) are considerably higher than those previously reported by other investigators (Barnard et al., 1974; Wranne and Woodson, 1973). Since the rats in the present study were younger than those in Barnard et al. (1974) and Wranne and Woodson's (1973) studies, it is possible that the elevation was due to the use of younger animals. Another explanation may be the different spacial orientations of the heart. When the rat is running on the horizontal as in the two former studies, the relationship between the heart and the body may be likened to the supine position in man, where stroke volume is at maximum due to maximal venous return (Edington and Edgerton, 1977) and hence heart rates may not be elevated to the same extent to meet the same cardiac outputs. On the vertical treadmill, the heart must work against the forces of gravity to move blood from the lower extremities and to the upper extremities. This may drop stroke volume and hence heart rates
must be elevated to meet the required cardiac output. In the present study, a decrease in maximal heart rate with aging was observed which is consistent with the findings of Barnard et al (1974). This observation is similar to that reported in man (Shepherd, 1968; Robinson, 1978).

Pre-exercise heart rates (Table 3) are similar to those reported by Barnard et al. (1974) and Wranne and Woodson (1973). During the second experimental week, the pre-exercise heart rates were significantly higher than during the eighth week, suggesting that at the younger age the animals were more excitable. This may be due in part to their being much more familiar with the handing and exercising after six weeks. However, pre-exercise heart rates were still elevated over the respective resting values in the eighth week. This type of anticipation to exercise has also been shown to increase the oxygen consumption (Shepherd and Gollnick, 1976) and rectal temperature (Gollnick and Ianuzzo, 1968) of rats. The resting heart rates (Table 3) obtained after the animals had rested for at least one-half hour are similar to those reported by other investigators (Tipton et al., 1966; Gollnick and Ianuzzo, 1968; Barnard et al., 1974). Basal heart rates (Table 3) were recorded during the light portion of the animal's daily cycle and only reported if the animal remained asleep. Basal heart rates decreased significantly from the second to eighth experimental week.

On a number of occasions, heart rates were recorded in the 700 to 900 beats/min. range (Appendix G). This tachycardia usually occurred when the animal became frightened with the heart rate
returning to normal within a few seconds. In man maximal exercise heart rates are the same as those elicited under heavy emotional stress. It may be that the maximal exercise heart rates reported here and elsewhere are not exceeded during exercise because cardiac output would not be augmented due to decreased filling. However, when using the rat in exercise and emotionally stressful environments, the heart rate may be elevated beyond the functional range and hence became less efficient. If this is the case, it certainly suggests that electric shock as a motivator may be counter productive for optimal cardiac function during exercise.

The heart rate response of the rat to various levels of power output (Fig. 10) is different than the heart rate response observed in man. Figure 10 illustrates the rapid response of the heart to various power outputs. Popovic et al. (1969) have reported that stroke volume does not increase with exercise in the rat. Therefore, it appears that any increase in cardiac output would come solely from an increase in heart rate. Hence, in order to elevate cardiac output heart rate is at or near maximal values at the low power outputs (Figure 10). Since oxygen consumption continues to rise at power output levels above the levels where heart rate has levelled off, the increased oxygen consumption must be achieved by enhanced extraction at the tissue level. Certainly the oxidative potential of rat skeletal muscle responds very quickly to aerobic conditioning programs (Holloszy et al., 1975). Table 4 shows the data obtained on 3 animals running at varying speeds and inclinations on a traditional horizontal treadmill. When speed was held constant and grade increased or vice versa

an increase in oxygen consumption was observed. For example, at $26.8 \mathrm{~m} / \mathrm{min}$. BL3 increased his oxygen consumption from $53.2 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. at $4^{\circ}$ grade to $68.7 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. at $10^{\circ}$ grade to a further increase of $78.2 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. at $16^{\circ}$ grade. This is in agreement with Wilson et al. (1978) and follows the same pattern that exists in man (Mathews and Fox, 1976). However, it is in disagreement with the work of Shepherd and Gollnick (1976) and does not support the findings of Taylor et al. (1972). Taylor et al (1972) have reported that in animals weighing less than 1 kg body weight, the metabolic cost of running is not significantly changed by the angle of the running surface. Shepherd and Gollnick (1976) support this finding with rats using a running wheel. Therefore, since both speed and grade on the horizontal contribute to the metabolic overload on the animal, comparisons between studies wherein different grades and speeds are used become tenuous. If standard elevations could be adapted it would certainly improve the generalizability of the results. The vertical treadmill used in this study offers a reasonable alternative and permits the calculation of physical work and power.

A closer approximation of the power output on the horizontal treadmill can be estimated from the oxygen consumption collected at a specific speed and grade. Assuming that a given power output on either the vertical or horizontal requires the same amount of oxygen consumption, then the individual regression lines found in Appendix E could be used to estimate power output on the horizontal treadmill, if oxygen consumption was known.

General Discussion

One of the most common changes that occurs with physical conditioning is a lower heart rate at submaximal power outputs (Astrand, 1977). A common practice of the majority of researchers conditioning rats is to progressively increase the speed of the treadmill and duration of the run during the first few weeks. The animals are then maintained at a selected speed and duration throughout the remainder of the experiment. It is presumed that during the first few weeks the rat undergoes a progressive overload. However, once the criterion speed is reached and no further increase takes place it is doubtful as to whether an optimal conditioning effect occurs. In an attempt to overcome this problem, the heart rates of rats conditioned in the present study were monitored. The heart rates were maintained in a range of 580 to 600 beats per minute by varying the speed. A progressive increase in the power output was observed throughout and was a function of increasing speed and/or increasing body weight (see Appendix H).

Another common practice when conditioning rats is to utilize a multi-animal treadmill and therefore condition five to ten rats at the same time. When heart rates are measured at a common speed and grade, they vary considerably as shown in Table 4. For example, the three rats measured at $26.8 \mathrm{~m} / \mathrm{min} ., 4^{\circ}$ grade demonstrated heart rates of 529,546 and 563 beats/min. This could lead to a large variation in the values of a number of dependent variables commonly investigated since the relative intensity at which each animal is being conditioned is different. However, at the faster speeds and/or greater inclines, the heart
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rates are more similar (Table 4). Since conditioning rats by heart rates is very time consuming and not practical if a large number is required, it is suggested that a grade of at least $10^{\circ}$ and a speed of approximately $30 \mathrm{~m} / \mathrm{min}$. be employed.

In the past, many investigators utilized changes in heart rate, muscle glycogen, etc., in an attempt to evaluate the strenuousness of the exercise since the measurement of oxygen consumption was difficult. Shepherd and Gollnick (1976) have suggested that the results from their study can be extrapolated to any type of running device where rats run at controlled speeds. They have assumed that an increase in grade does not increase metabolic rate as was discussed earlier. If their data is applied to the rats in the present study that were measured on the horizontal treadmill, then a gross over-estimation of their oxygen consumption would result. Therefore, it is suggested that the use of running speed to evaluate the strenuousness of the exercise bout is improper.

In man, the linear relationship between heart rate and oxygen consumption at submaximal power outputs allows the use of heart rates to estimate the strenuousness of exercise (Astrand, 1977). This is possible in man because of the $200 \%$ increase in heart rate from resting to maximal exercise allowing a wide range. However in the rat, heart rate increases only $50 \%$. Therefore at low power outputs the heart rate is at near maximal values, while oxygen consumption is still at submaximal levels. This suggests that it would be dubious to predict oxygen consumption in the rat from heart rate.

There are many similarities with regard to the rat and man
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such as the linear relationship between oxygen consumption and
power output, etc. However, there are a few dissimilarities such as
the rapid increase to maximum heart rate which render the rat as
less than an excellent experimental model for studying the systemic
adaptation to exercise. However, it is suggested that for the
study of peripheral adaptations, the rat remains a more than
adequate model since the increase in oxygen consumption would come
mainly from the extraction of oxygen at the tissue level and not
because of an increase in cardiac output.
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(2)

Conclusions

1. Body weight was lowest in the experimental group which was exposed to the greatest amount of total work. It appears that body weight changes are not only dependent upon the age and sex of the animal but also upon the duration, intensity and total work of the exercise and/or conditioning program.
2. The conditioning program failed to produce a significant difference in maximal oxygen consumption between the experimental groups in spite of substantially different amounts of chronic exercise. However, the slopes of the regression lines for the conditioned and non-conditioned groups were significantly different. This suggests that an adaptation in the conditioned group has occured.
3. A linear relationship exists between oxygen consumption (1/min) and power output.
4. Heart rate approaches maximal values at low power output levels. However, oxygen consumption continues to rise at power out put levels where heart rate has levelled off. This suggests that the increased oxygen consumption must be achieved by enhanced extraction at the tissue level.
5. The different slopes in conjunction with similar oxygen consumption values for the different groups suggest that the rat may not be an ideal model for man when studying exercise adaptations in the oxygen transport system. However, it is suggested that for the study of peripheral adaptations, the rat remains a more than adequate model since the increase in oxygen consumption would come
from the extraction of oxygen at the tissue level and not because of an increase in cardiac output.

Recommendations

1. There is a need for further research to determine if a mathematical relationship exists between different mechanical and metabolic power outputs on the vertical and horizontal treadmills.
2. Future investigation might be aimed at determining the aerobic - anaerobic components of both conditioned and non-conditioned rats at varying power outputs.
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APPENDIX A

REVIEW OF RELEVANT LITERATURE
(2)

REVIEW OF RELEVANT IITERATURE

During the past ten years, researchers have shown a substantial number of physiological changes to occur in the rat with conditioning. The conditioning protocols that have been used are markedly different, thereby making comparisons between studies extremely difficult. In the present study, a vertical treadmill was utilized which allows the power output performed by the rat to be measured. Along with the measurement of metabolic work and the heart rate responses to various power outputs, the ability to Quantify the conditioning stimulus was enhanced.

MODES OF CONDITIONING

A variety of methods have been used by investigators in an attempt to study exercise-related phenomena in small animals. The most common methods used are running animals on a motor-driven treadmill, running wheel, exercise drum or exposing the animal to swimming with an additional weight attached to the tail or thorax.

Until recently, motor-driven treadmills were homemade
(Koniski, 1960; Critz and Merrick, 1962; Andrews, 1965; Brannon et al., 1968). The grade and speed of the treadmill could be altered to increase the power output of the animal. Electrical stimulation was employed as a motivational device for teaching animals to run. Some control over the power outputs was provided. However, there are a number of problems associated with conditioning animals on a motor-driven treadmill. If the treadmill accomodates several animals at a time then the running speed must be adjusted to the

slowest animal (Wells and Heusner, 1971). The power output performed by each animal is difficult to assess when they are conditioned in groups (Hardin, 1965). All animals can not be induced to train on a motor-driven treadmill every day and some refuse to run thereby being dragged or rolled on the treadmill surface with the possibility of injury occuring (Thomas and Miller, 1958; Andrews, 1965; Hanson et al., 1969). Frequently, animals will be bruised and cut in the area of the lower limb (Reardon, 1975) or develop sore feet from the surface of the treadmill, thereby missing sessions or being eliminated from the study (Hardin, 1965).

Wells and Heusner (1971) developed a controlled running
wheel for small animals in an attempt to eliminate many of the above problems that occur with motor-driven treadmills. Groups of animals can be conditioned simultaneously with each animal allowed to respond individually to their running program. The speed and duration of running can be regulated, therefore continuous or intermittent conditioning programs are possible. A major drawback with this type of conditioning mode is that the power output is difficult to measure, since the animals do not always run in the same position in the wheel (Hardin, 1965).

Jenness and Byrd (1974) have recently designed a "live-in" exercise drum unit capable of housing and conditioning forty animals simultaneously. Handling of animals is eliminated, however animals can not be conditioned individually. The reported speed range is from 7.32 - 43.89 meters per minute, thus high intensity anaerobic conditioning would not be possible. The authors also
reported sore paw problems while conditioning a group for five days at 7.32 meters $/ \mathrm{min}$. for fifteen minutes each day. Calculation of the power output each animal performed would also be difficult with this apparatus.

Swimming has been a popular form of exercise employed by many laboratories. The equipment necessary is relatively inexpensive and simple. The laboratory rat requires no training because of their natural swimming ability. However, a number of factors can influence the swimming times of animals.

A number of investigators have shown marked reductions in swimming times of rats forced to swim in water temperatures different from body temperature (Tan et al., 1954; Baker and Horvath, 1964a; Beaton and Feleki, 1967).

Baker and Horvath (1964b) have reported that the swimming style of all animals is not the same. Some animals never allow their nostrils to drop below the surface of the water, whereas other animals sank for several seconds and then rose to the surface and swam for several seconds. A few rats would sink to the bottom of the tank, remain in a vertical position and exhale through their nostrils for as long as 20 seconds. This procedure was followed by a rapid ascent to the surface where swimming would take place for several seconds and then the sinking process was repeated.

In an attempt to eliminate the various styles of swimming by standardizing the power output, several methods of weighing rats have been employed (Hardin, 1865; 1968; Dawson and Horvath,

1970). Another source of variability in swimming is the possibility of trapped air bubbles in the fur which increases the buoyancy of the animals (Dawson and Horvath, 1970). Yevick et al. (1969) have reported an increase in chronic murine pneumonia with swimming rats as compared to the sedentary controls. This would imply that longitudinal studies in which swimming is used may not be advisable. In addition to the above factors that can affect swimming performance, the quantification of power output while swimming is not possible.

OXYGEN CONSUMPTION STUDIES (RAT)

Relatively little is known about the oxygen consumption of the rat during maximal exercise. Shepherd and Gollnick (1976) adapted a running wheel to determine oxygen consumption at a number of different speeds ( $16-67 \mathrm{~m} / \mathrm{min}$.$) during a four to five$ week period. Their results demonstrated that oxygen consumption increased linearly with work intensity (running speed) up to maximum. A maximum oxygen consumption of $95.1 \pm 1.4 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. was found to occur at a running speed of $49.5 \mathrm{~m} / \mathrm{min}$. The authors suggest that their results can be extrapolated to any type of running device where rats run at controlled speeds.

Recently, Brooks and White (1977) reported submaximal values of oxygen consumption around $80 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. in rats exercising at $43.1 \mathrm{~m} / \mathrm{min}$. on a $11.3^{\circ}$ slope. According to Shepherd and Gollnick's (1976) data a speed of $43.1 \mathrm{~m} / \mathrm{min}$. produces an oxygen consumption of approximately $88 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. The $8 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. difference could possibily be due to a younger rat used in the Brooks and White study
or a difference in the training modes used. The mean weight of their rats was 222 grams, whereas the rats used in the Shepherd and Gollnick study had a mean weight of 373 grams.

Taylor et al. (1970) developed a regression equation to predict oxygen consumption from running speed. Prediction of an oxygen consumption from their equation yields lower values than the equation developed by Shepherd and Gollnick (1976).

Wilson et al.(1978) have recently reported maximal values of oxygen consumption for untrained male and female (Sprague-Dawley and Okamato-Aoki strains)rats. A graded treadmill test produced maximal oxygen consumptions of $91 \pm 11$ and $87 \pm 11 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. for male and female rats respectively. Maximal oxygen consumption was elicited at a treadmill speed of $30.3 \mathrm{~m} / \mathrm{min}$. with a $15^{\circ}$ inclination. The authors have shown an increase in oxygen consumption with a progressive grade and speed.

Oxygen consumption in the swimming rat has been studied by a number of investigators (Baker and Horvath, 1964b; Kratzing and Mark, 1967; McArdle, 1967; Dawson and Horvath, 1970). Maximum values have been reported to range from 55 to $80 \mathrm{ml} / \mathrm{kg} / \mathrm{min}$. Comparisons between the studies becomes difficult because of the various methodologies utilized by each laboratory.

McArdle (1967) suggests that the addition of weight does not increase the oxygen consumption in many animals, but actually reduces the ability of the animal to exchange gases at the surface. Therefore, swimming may lend itself to submaximal exercise programs but is less suited to programs of near maximal intensity (Hardin, 1965; McArdle, 1967).

EVALUATION OF WORK CAPACITY

Oscai and Molé (1975) suggest that the work capacity of animals can be evaluated by subjecting them to a run or swim to exhaustion and then compare the results of the conditioned versus the control animals. The animal may be considered exhausted when it can no longer right itself after being placed on its back (Oscai and Mole, 1975) or if the animal sits at the back of the treadmill for a period of more than 10 seconds, despite being subjected to electric shock (Jobin, 1977).

A graded treadmill test for rats has been developed by Wranne and Woodson (1973) for determination of maximal work performance. The speed and inclination are progressively increased from $26.8 \mathrm{~m} / \mathrm{min}$. at $12.5^{\circ}$ grade to $34.8 \mathrm{~m} / \mathrm{min}$. at a $27^{\circ}$ grade or until exhaustion occurs. A linear increase in heart rate with an increasing power output occurred from a average of $412 \pm 20$ beats/ min. pre-exercise to an average of $615 \pm 12$ beats/min. at maximum. Arterial lactate also increased progressively from rest to maximum. The authors suggest that the test is suited to detect changes in blood oxygen transport.

OXYGEN CONSUMPTION STUDIES (MAN)
The determination of maximal oxygen consumption in man has received considerable attention in the literature. A variety of continuous (Balke, 1959; Cureton, 1969; Kamon and Pandolf, 1972b; Tolson and Ismail, 1970) and discontinuous (Taylor et al., 1955; Mitchell et al., 1958; Astrand et al., 1959; Macnab et al., 1969) tests have been developed to elicit maximal oxygen consumption

utilizing the treadmill, bicycle ergometer and laddermill.
It was determined by Taylor et al. (1955) that increasing the speed and keeping the grade constant was not as effective for obtaining an asymptote as increasing the grade with the speed held constant. Shepherd and Gollnick (1976) reported that the increase in grade was not as important as increasing the speed in rats which is contrary to what has been observed in man.

The precise power ouptut can be obtained on the bicycle ergometer and laddermill, whereas, the power output of uphill running is confounded by a horizontal component (Kamon and Pandolf, 1972b).

Kamon (1972a) reported a linear relationship between oxygen consumption and power output on both the laddermill and bicycle ergometer. Laddermill climbing was found to elicit maximal oxygen consumption values similar to uphill running but higher than cycling (Kamon and Pandolf, 1972b).

Because of the complexity of the direct measurement of oxygen consumption as well as the physical demand made upon the individual, a number of indirect methods of predicting maximal oxygen consumption have been developed. The indirect tests assume a linear relationship between heart rate and oxygen consumption and therefore extrapolation to a predicted maximum heart rate will yield a prediction of maximal oxygen consumption (Anderson et al., 1971). One of the most popular indirect tests was developed by Astrand and Ryhming (1954). A nomogram is used to compute maximal oxygen consumption from the heart rate after a submaximal
bicycle ergometer ride. When this test is compared to a maximum bicycle test predicted maximal oxygen consumption is higher (Glassford et al., 1965). The reason for this may be due to the fact that the submaximal test to predict maximal oxygen consumption is not as fatiguing on the leg muscles as the maximal test.

Hermiston and Faulkner (1971) used a stepwise multipleregression technique to develop equations for the prediction of maximal oxygen consumption of physically active and inactive men. For example, the equation for active men takes into account the age of the subject, fat free weight, heart rate, fraction of carbon dioxide in expired gas, tidal volume at submaximal work level and respiratory exchange ratio. The correlation coefficient between the observed and predicted oxygen consumption was found to be 0.90. Other investigators (Mastopaolo, 1970; Metz and Alexander, 1971; Jessup et al., 1974) have improved upon the predictive validity of simple regression models by including a wide variety of respiratory and cardiovascular variables.

For the prediction of maximal oxygen consumption in man, a considerable body of knowledge has accumulated in the literature over the years. However, our knowledge of the physical work capacity of the rat is extremely limited. There has been a scarcity of material in the literature concerning the rat, therefore researchers have not been able to quantify their training programs. If we are to continue to use the rat to study the effects of exercise then the relative intensity of their conditioning programs must be known.

APPENDIX B

RAW DATA







Appendix B-2 Raw data for maximal oxygen consumption test on the vertical treadmill

| Group | Animal Number | Weight (grams) | Power Output (watts) | Heart Rate (b/min) | $\begin{gathered} \stackrel{\dot{V} O_{2}}{(1 / \mathrm{min})} \end{gathered}$ | $\begin{gathered} \dot{\mathrm{V} O} \\ (\mathrm{ml} / \mathrm{kg} / \\ \mathrm{min}) \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| SC | GRR | 370.7 | Pre-exercise | 445 | 0.0150 | 40.46 |
|  |  |  | 0.519 | 558 | 0.0235 | 63.39 |
|  |  |  | 0.821 | 593 | 0.0269 | 72.57 |
|  |  |  | 0.958 | 600 | 0.0301 | 81.20 |
| SC | GRP | 396.9 | Pre-exercise | 420 | 0.0121 | 30.49 |
|  |  |  | 0.556 | 563 | 0.0256 | 64.50 |
|  |  |  | 0.691 | 579 | 0.0296 | 74.58 |
|  |  |  | 0.804 | 600 | 0.0321 | 80.88 |
|  |  |  | 0.947 | 610 | 0.0343 | 86.42 |
|  |  |  | 1.108 | 600 | 0.0331 | 83.44 |
| SC | RBK | 347.9 | Pre-exercise | 400 | 0.0104 | 29.89 |
|  |  |  | 0.487 | 558 | 0.0200 | 57.49 |
|  |  |  | 0.606 | 585 | 0.0211 | 60.65 |
|  |  |  | 0.771 | 600 | 0.0273 | 78.47 |
|  |  |  | 0.899 | 600 | 0.0280 | 80.48 |
| SC | RBL | 360.4 | Pre-exercise | - | 0.0111 | 30.80 |
|  |  |  | 0.505 | - | 0.0255 | 70.45 |
|  |  |  | 0.628 | - | 0.0270 | 74.92 |
|  |  |  | 0.730 | - | 0.0299 | 82.96 |
|  |  |  | 0.860 | - | 0.0327 | 90.73 |
|  |  |  | 0.931 | - | 0.0302 | 83.80 |



Appendix B-2 continued

| Group | Animal Number | Weight (grams) | Power Output (watts) | Heart Rate (b/min) | $\begin{gathered} \stackrel{\mathrm{VO}}{2}^{(1 / \mathrm{min})} \end{gathered}$ | $\begin{aligned} & \dot{\mathrm{VO}}_{2} \\ & (\mathrm{ml} / \mathrm{kg} / \\ & \mathrm{min}) \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| SC | GBK | 397.2 | Pre-exercise | - | 0.0114 | 28.70 |
|  |  |  | 0.557 | - | 0.0248 | 62.44 |
|  |  |  | 0.692 | - | 0.0259 | 65.21 |
|  |  |  | 0.745 | - | 0.0279 | 70.24 |
|  |  |  | 0.805 | - | 0.0270 | 68.07 |
| SC | RPL | 347.2 | Pre-exercise | - | 0.0121 | 34.85 |
|  |  |  | 0.411 | - | 0.0178 | 51.27 |
|  |  |  | 0.549 | - | 0.0183 | 52.71 |
|  |  |  | 0.769 | - | 0.0212 | 61.06 |
|  |  |  | 0.897 | - | 0.0249 | 71.72 |
| SC | GRB | 358.6 | Pre-exercise | - | 0.0109 | 30.40 |
|  |  |  | 0.424 | - | 0.0211 | 58.84 |
|  |  |  | 0.567 | - | 0.0237 | 66.09 |
|  |  |  | 0.727 | - | 0.0246 | 68.60 |
|  |  |  | 0.794 | - | 0.0234 | 65.25 |
| SC | BLB | 426.5 | Pre-exercise | 390 | 0.0118 | 27.67 |
|  |  |  | 0.505 | 585 | 0.0285 | 66.82 |
|  |  |  | 0.674 | 600 | 0.0315 | 73.86 |
|  |  |  | 0.864 | 610 | 0.0335 | 78.55 |
|  |  |  | 0.945 | 610 | 0.0339 | 79.48 |
|  |  |  | 1.018 | 600 | 0.0332 | 77.84 |



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Appendix B-2 continued
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| Group | Animal Number | Weight <br> (grams) | Power Output (watts) | Heart Rate (b/min) | $\begin{gathered} \stackrel{\mathrm{VO}}{2}^{(1 / \mathrm{min})} \end{gathered}$ | $\begin{aligned} & \mathrm{VO}_{2} \\ & (\mathrm{ml} 7 \mathrm{~kg} / \\ & \mathrm{min}) \end{aligned}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| E1 | BLI | 377.8 | Pre-exercise | 364 | 0.0124 | 33.17 |
|  |  |  | 0.441 | 585 | 0.0200 | 53.65 |
|  |  |  | 0.559 | 581 | 0.0225 | 60.22 |
|  |  |  | 0.589 | 590 | 0.0262 | 70.28 |
|  |  |  | 0.699 | 600 | 0.0278 | 72.70 |
|  |  |  | 0.755 | 600 | 0.0265 | 71.08 |
| E1 | BKI | 307.2 | Pre-exercise | 393 | 0.0139 | 45.27 |
|  |  |  | 0.364 | 578 | 0.0162 | 52.73 |
|  |  |  | 0.486 | 580 | 0.0215 | 69.99 |
|  |  |  | 0.576 | 600 | 0.0232 | 75.52 |
|  |  |  | 0.681 | 581 | 0.0221 | 72.04 |
| El | GRI | 331.5 | Pre-exercise | 375 | 0.0142 | 42.98 |
|  |  |  | 0.392 | 571 | 0.0169 | 50.98 |
|  |  |  | 0.459 | 578 | 0.0199 | 60.03 |
|  |  |  | 0.524 | 580 | 0.0222 | 66.97 |
|  |  |  | 0.621 | 571 | 0.0283 | 85.37 |
|  |  |  | 0.734 | 571 | 0.0254 | 76.62 |
| El | PBK | 337.9 | Pre-exercise | 368 | 0.0128 | 37.88 |
|  |  |  | 0.473 | 541 | 0.0201 | 59.49 |
|  |  |  | 0.589 | 565 | 0.0243 | 71.92 |
|  |  |  | 0.685 | 571 | 0.0259 | 76.65 |
|  |  |  | 0.749 | 571 | 0.0245 | 72.51 |



Appendix B-2 continued

| Group | Animal <br> Number | Weight (grams) | Power Output (watts) | Heart Rate (b/min) | $\stackrel{\text { ㅂㅇ }}{2}_{(1, \mathrm{~m} \mathrm{n})}$ | $\begin{gathered} \stackrel{\mathrm{VO}}{2} \\ (\mathrm{ml} / \mathrm{kg} / \\ \mathrm{min}) \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| E1 | GBB | 339.8 | Pre-exercise | 416 | 0.0150 | 44.14 |
|  |  |  | 0.476 | 545 | 0.0183 | 53.86 |
|  |  |  | 0.592 | 554 | 0.0196 | 57.68 |
|  |  |  | 0.637 | 577 | 0.0215 | 63.27 |
|  |  |  | 0.688 | 580 | 0.0225 | 66.22 |
|  |  |  | 0.753 | 600 | 0.0210 | 61.79 |
|  |  |  | 0.811 | 590 | 0.0190 | 55.97 |
| E2 | PR4 | 363.2 | Pre-exercise | 436 | 0.0183 | 50.27 |
|  |  |  | 0.430 | 600 | 0.0216 | 59.47 |
|  |  |  | 0.574 | 600 | 0.0225 | 61.92 |
|  |  |  | 0.681 | 610 | 0.0240 | 66.08 |
|  |  |  | 0.736 | 600 | 0.0254 | 69.93 |
|  |  |  | 0.805 | 590 | 0.0238 | 65.53 |
| E2 | BK4 | 368.4 | Pre-exercise | 420 | 0.0101 | 27.42 |
|  |  |  | 0.516 | 566 | 0.0191 | 51.85 |
|  |  |  | 0.642 | 581 | 0.0225 | 61.07 |
|  |  |  | 0.746 | 600 | 0.0269 | 73.02 |
|  |  |  | 0.816 | 600 | 0.0241 | 65.42 |
| E2 | GR4 | 400.1 | Pre-exercise | 393 | 0.0120 | 29.99 |
|  |  |  | 0.474 | 581 | 0.0209 | 52.24 |
|  |  |  | 0.633 | 600 | 0.0245 | 61.24 |
|  |  |  | 0.750 | 610 | 0.0261 | 65.23 |
|  |  |  | 0.811 | 600 | 0.0251 | 62.73 |




E2
410.1 Pre-exercise

413
0.0098
23.90

585
0.018545 .11

590
$0.0224 \quad 54.62$
605
$0.0264 \quad 64.38$
600
$0.0253 \quad 61.69$

Cl PR3
$313.1 \quad$ Pre-exercise 0.371
421
$0.0148 \quad 47.21$
0.015248 .55
$0.0197 \quad 62.92$
$0.0234 \quad 74.74$
$0.0260 \quad 83.04$
600
0.024176 .81

C1 PRR 313.8 Pre-exercise
436
$0.0099 \quad 31.62$
0.371

571
$0.0169 \quad 53.86$
0.496

600
$0.0206 \quad 65.65$
0.587

600
$0.0264 \quad 84.13$
0.634600
$0.0241 \quad 76.80$
$\mathrm{Cl} \quad \mathrm{BK} 3 \quad 306.6$
Pre-exercise
0.363
0.428
0.534
0.679
0.792

390
$0.0110 \quad 35.82$
$0.0186 \quad 60.67$
571
$0.0204 \quad 66.54$
590
$0.0254 \quad 82.84$
0.679571
$0.0216 \quad 70.46$
0.792

568
$0.0214 \quad 69.80$


Appendix B-2 Continued

| Group | Animal Number | Weight <br> (grams) | Power Output (watts) | Heart Rate (b/min) | $\stackrel{\mathrm{VO}}{2}^{(1 / \mathrm{min})}$ | $\begin{gathered} \stackrel{\dot{\mathrm{VO}}}{(\mathrm{ml} / \mathrm{kg} /} \underset{\mathrm{min})}{ } \end{gathered}$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Cl | BL3 | 241.1 | Pre-exercise | 414 | 0.0094 | 38.60 |
|  |  |  | 0.286 | 545 | 0.0137 | 56.59 |
|  |  |  | 0.341 | 564 | 0.0151 | 62.37 |
|  |  |  | 0.383 | 564 | 0.0172 | 71.05 |
|  |  |  | 0.424 | 571 | 0.0184 | 76.00 |
|  |  |  | 0.536 | 581 | 0.0211 | 87.15 |
|  |  |  | 0.676 | 545 | 0.0195 | 80.55 |
| C2 | GR2 | 366.8 | Pre-exercise | 413 | 0.0140 | 37.99 |
|  |  |  | 0.434 | 565 | 0.0258 | 70.34 |
|  |  |  | 0.639 | 578 | 0.0276 | 75.25 |
|  |  |  | 0.743 | 580 | 0.0307 | 83.70 |
|  |  |  | 0.873 | 554 | 0.0281 | 76.81 |
| C2 | BL2 | 303.2 | Pre-exercise | 364 | 0.0098 | 32.32 |
|  |  |  | 0.359 | 565 | 0.0162 | 53.43 |
|  |  |  | 0.479 | 576 | 0.0191 | 63.02 |
|  |  |  | 0.528 | 581 | 0.0218 | 71.92 |
|  |  |  | 0.568 | 585 | 0.0252 | 83.14 |
|  |  |  | 0.614 | 585 | 0.0248 | 81.79 |
| C2 | BK2 | 301.8 | Pre-exercise | 393 | 0.0102 | 33.80 |
|  |  |  | 0.357 | 545 | 0.0198 | 65.61 |
|  |  |  | 0.423 | 576 | 0.0214 | 70.91 |
|  |  |  | 0.477 | 590 | 0.0223 | 73.89 |
|  |  |  | 0.526 | 590 | 0.0252 | 83.50 |
|  |  |  | 0.611 | 581 | 0.0243 | 80.52 |



Appendix B-3 Raw data for heart rates during the 2 nd. week

| Group | Basal | Resting <br> in cage | Pre-exercise | Maximum Exercise |
| :---: | :---: | :---: | :---: | :---: |
| SC | - | - | 462 | 655 |
|  | - | - | 514 | 679 |
|  | - | - | 545 | 643 |
|  | - | - | 480 | 632 |
|  | 379 | 405 | 467 | 658 |
|  | - | - | 500 | 643 |
|  | - | - | 500 | 632 |
|  | - | - | 525 | 655 |
| E1 | 384 | 412 | 494 | 655 |
|  | - | - | 505 | 655 |
|  | 424 | 486 | 525 | 624 |
|  | - | - | 494 | 658 |
| E2 | 408 | 500 | 525 | 667 |
|  | - | - | 500 | 655 |
|  | - | - | 494 | 632 |
|  | - | - | 500 | 624 |
|  | - | - | - | 667 |
| Cl | - | - | 480 | 647 |
|  | 405 | 384 | 480 | 658 |
|  | - | - | $514$ | 658 |
|  | - | - | 545 | 643 |
| C2 | 444 | 494 | 500 | 621 |
|  | - | - | 494 | 667 |
|  | - | - | 514 | 632 |
|  | - | - | 545 | 624 |
|  | - | - | 494 | 643 |

Appendix B-4 Raw data for heart rates during the 8 th. week

| Group | Basal | Resting <br> in cage | Pre-exercise | Maximum Heart Rate |
| :---: | :---: | :---: | :---: | :---: |
| SC | - | - | 445 | 600 |
|  | 305 | 380 | 420 | 610 |
|  | - | - | 400 | 600 |
|  | - | - | - | - |
|  | - | - | - | - |
|  | - | - | - | - |
|  | - | - | - | - |
|  | - | - | 390 | 610 |
| El | 269 | 335 | 364 | 600 |
|  | - | - | 393 | 600 |
|  | - | - | 375 | 580 |
|  | 315 | 333 | 368 | 571 |
|  | - | - | 416 | 600 |
| E2 | - | - | 436 | 610 |
|  | - | - | $420$ | 600 |
|  | - | - | $393$ | $610$ |
|  | - | - | $413$ |  |
| Cl | - | - | 421 | 615 |
|  | - | - | 436 | 600 |
|  | $305$ | - | $390$ | $593$ |
|  | $307$ | 369 | 414 | 581 |
| C2 | - | - | 413 | 580 |
|  | - | - | 364 | 585 |
|  | - | - | 393 | 590 |



APPENDIX C
DESCRIPTION OF VERTICAL TREADMILL


Shaft and 2" Idler Sprockets

3/8" Roller Chains Connected By 3/16" Rods, 3/4" Pitch


## RAT FRCDIVETER



APPENDIX D
TRAINING PROGRSSION


Appendix D-l Training progression for groups El, E2 and SC

| Day | Time | Speed <br> $(\mathrm{m} / \mathrm{min})$ | Rep. | Total on <br> (min:sec) | Total off <br> $(\mathrm{min}: \mathrm{sec})$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | AM | 5.2 | 1 | $00: 30$ | - |
| 2 | AM | 5.2 | 2 | $01: 00$ | $00: 30$ |
| 3 | AM | 5.2 | 3 | $01: 30$ | $00: 30$ |
| 5 | AM | 5.2 | 3 | $02: 15$ | $00: 30$ |
| 6 | AM | 5.2 | 3 | $03: 00$ | $00: 30$ |

From this point on, the animals in group SC were trained for a duration of 1 - 2 min . once a week, whereas animals in groups El and E2 were exercised at a heart rate range of 580 to 610 beats/min.

| 12 | AM | 1 | 03:00 | - |
| :---: | :---: | :---: | :---: | :---: |
|  | PM | 1 | 03:00 | - |
| 16 | AM | 1 | 01:00 | - |
|  | PM | 1 | 01:00 | - |
| 19 | AM | 1 | 04:00 | - |
|  | PM | 1 | 04:00 | - |
| 23 | AM | 1 | 01:00 | - |
|  | PM | 1 | 01:00 | - |
| 26 | AM | 1 | 05:00 | - |
|  | PM | 1 | 05:00 | - |
| 30 | AM | 1 | 01:00 | - |
|  | PM | 1 | 01:00 | - |
| 33 | AM | 1 | 06:00 | - |
|  | PM | 1 | 06:00 | - |
| 37 | AM | 1 | 01:00 | - |
|  | PM | 1 | 01:00 | - |
| 40 | AM | 1 | 07:00 | - |
|  | PM | 1 | 07:00 | - |
| 44 | AM | 1 | 01:00 | - |
|  | PM | 1 | 01:00 | - |
| 47 | AM | 1 | 08:00 | - |
|  | PM | 1 | 08:00 | - |
| 51 | AM | 1 | 01:00 | - |
|  | PM | 1 | 01:00 | - |

Appendix D-2 Training progression for groups Cl and C2

| Day | Time | Speed <br> $(\mathrm{m} / \mathrm{min})$ | Rep. | Total on <br> $(\mathrm{min}: \mathrm{sec})$ | Total off <br> $(\mathrm{min}: \mathrm{sec})$ |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 1 | AM | 5.2 | 1 | $00: 30$ | - |
| 2 | AM | 5.2 | 2 | $01: 00$ | $00: 30$ |
| 3 | AM | 5.2 | 3 | $01: 00$ | $00: 30$ |
| 5 | AM | 5.2 | 3 | $02: 15$ | $00: 30$ |
| 6 | AM | 5.2 | 3 | $03: 00$ | $00: 30$ |

From this point on, the animals in groups Cl and C2 were conditioned at a heart rate range of 590 to 610 beats/min.

| 8 | AM | 1 | 03:00 | - |
| :---: | :---: | :---: | :---: | :---: |
|  | PM | 1 | 03:00 | - |
| 10 | AM | 1 | 04:00 | - |
|  | PM | 1 | 04:00 | - |
| 11 | AM | 1 | 05:00 | - |
|  | PM | 1 | 05:00 | - |
| 14 | AM | 1 | 06:00 | - |
|  | PM | 1 | 06:00 | - |
| 15 | AM | 1 | 07:00 | - |
|  | PM | 1 | 07:00 | - |
| 17 | AM | 1 | 08:00 | - |
|  | PM | 1 | 08:00 | - |
| 18 | AM | 1 | 09:00 | - |
|  | PM | 1 | 09:00 | - |
| 21 | AM | 1 | 10:00 | - |
|  | PM | 1 | 10:00 | - |
| 22 | AM | 1 | 11:00 | - |
|  | PM | 1 | 11:00 | - |
| 24 | AM | 1 | 12:00 | - |
|  | PM | 1 | 12:00 | - |
| 25 | AM | 1 | 13:00 | - |
|  | PM | 1 | 13:00 | - |
| 28 | AM | 1 | 14:00 | - |
|  | PM | 1 | 14:00 | - |
| 29 | AM | 1 | 15:00 | - |
|  | PM | 1 | 15:00 | - |
| 31 | AM | 1 | 15:00 | - |
|  | PM | 1 | 15:00 | - |
| 32 | AM | 1 | 15:00 | - |
|  | PM | 1 | 15:00 | - |
| 35 | AM | 1 | 15:00 | - |
|  | PM | 1 | 15:00 | - |

Appendix D-2 continued

| Day | Time | $\begin{aligned} & \text { Speed } \\ & (\mathrm{m} / \mathrm{min}) \end{aligned}$ | Rep. | Total on (min:sec) | Total off (min:sec) |
| :---: | :---: | :---: | :---: | :---: | :---: |
| 36 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 38 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 39 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 42 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 43 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 45 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 46 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 49 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |
| 50 | AM |  | 1 | 15:00 | - |
|  | PM |  | 1 | 15:00 | - |

APPENDIX E

INDIVIDUAL REGRESSION LINES
 the sedentary control group (SC)





APPENDIX F

SUMMARY OF STATISTICAL TESTS

```
Appendix F-1, a Summary of the analysis of variance for body
weights of sedentary control, exercised and
conditioned rats at the time of arrival in
the laboratory
```

| Source of <br> Variation | DF | SS | MS | F |  |
| :--- | ---: | ---: | ---: | ---: | ---: |
| Groups | 4 | 401.63 | 100.41 | 1.61 | $>0.05$ |
| Error | 25 | 1555.62 | 62.21 |  |  |

$F_{0.05}(4,25) 2.76$

Appendix $F-1, b$ Summary of the analysis of variance for body weights of sedentary control, exercised and conditioned rats at the termination of the study

| Source of <br> Variation | DF | SS | MS | $F$ | $P$ |
| :--- | :---: | :---: | :---: | :---: | :---: | :---: |
| Group | 4 | 23134.4 | 5783.5 | 5.81 | $<0.01$ |
| Error | 19 | 18907.1 | 995.1 |  |  |

[^2]Appendix $F-1, c$ Newman-Keuls comparison between ordered means for body weight of sedentary control, exercised and conditioned rats at the end of the study

|  | C1 | C2 | E1 | SC | E2 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Means | 307.8 | 330.5 | 347.2 | 384.7 | 386.0 |
| 307.8 | 0.0 | 22.7** | 39.4** | 76.9** | 78.2** |
| 330.5 |  | 0.0 | 16.7* | 54.2** | 55.5** |
| 347.2 |  |  | 0.0 | $37.5 * *$ | 38.3** |
| 384.7 |  |  |  | 0.0 | 1.3 |
| 386.0 |  |  |  |  | 0.0 |
| $\mathrm{R}=$ |  | 2 | 3 | 4 | 5 |

The Multiplier is 3.7969
** $\mathrm{p}<0.01$

* $p<0.05$


```
Appendix F-2, a Summary of the analysis of variance for maximal
oxygen consumption (ml/kg/min) of sedentary
control, exercised and conditioned rats
```

| Source of <br> Variation | DF | SS | MS | $F$ | $P$ |
| :---: | :---: | :---: | :---: | :---: | :---: | :---: |
| Groups | 4 | 673.94 | 168.49 | 4.63 | $<0.01$ |
| Error | 19 | 690.72 | 36.35 |  |  |

```
Fo.01 (4, 19) 4.50
```

Appendix F-2, b Newman-Keuls comparison between ordered means for maximal oxygen consumption ( $\mathrm{ml} / \mathrm{kg} / \mathrm{min}$.) of sedentary control, exercised and conditioned rats

|  | E2 | E1 | SC | C1 | C2 |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Means | 68.14 | 75.29 | 78.61 | 83.45 | 84.29 |
| 68.14 | 0.00 | 7.15 | $10.47 *$ | $15.31 * *$ | $16.15 * *$ |
| 75.29 |  | 0.00 | 3.32 | 8.16 | 9.00 |
| 78.61 |  | 0.00 | 4.84 | 5.68 |  |
| 83.45 |  |  |  | 0.00 | 0.84 |
| 84.29 | 2 | 3 | 4 | 0.00 |  |
| $R=$ |  |  |  |  | 5 |

The Multiplier is 2.9021
** $\mathrm{p}<0.01$

* $\mathrm{P}<0.05$

Appendix $F-3$ Summary of the Analysis of variance for preexercise oxygen consumptions of sedentary control, exercised and conditioned rats

| Source of <br> Variation | DF | SS | MS |  |  |
| :--- | :--- | :--- | :--- | :--- | :--- |
| Groups | 4 | 311.76 | 77.94 | 1.87 | $>0.05$ |
| Error | 19 | 790.41 | 41.60 |  |  |

$\mathrm{F}_{0.05}(4,19) 2.90$

Appendix $F-4$ Summary of the analysis of variance for maximum heart rate of sedentary control, exercised and conditioned rats during the 8 th experimental week

| Source of <br> Variation | DF | SS | MS |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Groups | 4 | 1261.70 | 315.43 | 2.99 | $>0.05$ |
| Error | 15 | 1584.30 | 105.62 |  |  |

$F_{0.05}(4,15) 3.06$

```
Appendix F-5 Summary of the analysis of varianse for pre-exercise
    heart rate of sedentary control, exercised and
    conditioned rats during the 8th experimental week
```

| Source of <br> Variation | DF | SS | MS | P |  |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Groups | 4 | 4153.90 | 1038.48 | 2.27 | $>0.05$ |
| Error | 15 | 6877.30 | 458.49 |  |  |

[^3]

Appendix $F-6$ Summary of the analysis of variance for maximum heart rate of sedentary control, exercised and conditioned rats during the 2nd experimental week

| Source of <br> Variation | DF | SS | MS | P |  |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Groups | 4 | 619.97 | 154.99 | 0.58 | $>0.05$ |
| Error | 21 | 5634.07 | 268.29 |  |  |

$F_{0.05}(4,21) 2.84$
(1)

```
Appendix F-7 Summary of the analysis of variance for pre-exercise
    Heart rate of sedentary control, exercised and con-
    ditioned rats during the 2nd experimental week
```

| Source of |  |  |  |  |  |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Variation | DF | SS | MS | F |  |
| Groups | 4 | 340.79 | 85.20 | 0.14 | $>0.05$ |
| Error | 21 | 11766.58 | 560.31 |  |  |

$$
F_{0.05}(4,20) 2.87
$$




```
Appendix F-9 Summary of analysis of variance for maximal
    oxygen consumption (l/min) of sedentary control,
    exercised and conditioned rats
```

| Source of <br> Variation | DF | SS | MS | F |  |
| :--- | :--- | :--- | :--- | :--- | :--- |
| Groups | 4 | 0.00009 | 0.00002 | 2.26 | $>0.05$ |
| Error | 19 | 0.00017 | 0.000009 |  |  |

F0.05 (4,19) 2.90

APPENDIX G

ECG TRACINGS

Appendix $G$ An example of the ECG tracing for pre-exercise, submaximal exercise and paroxysmal tachycardia the rat (paper speed at $25 \mathrm{~mm} / \mathrm{sec}$ ).


Pre-exercise - 450 beats/min.


Submaximal exercise - 578 beats/min.


APPENDIX H

EXAMPLE OF PROGRESSIVE INCREASE IN POWER OUTPUT AT 580-600 B/M.

```
Appendix H Example of the increase in power output maintaining
    heart rate in the 580-600 b/min. range for BL3
```

| Experimental Day | $\begin{aligned} & \text { Speed } \\ & (\mathrm{m} / \mathrm{sec}) \end{aligned}$ | Weight (grams) | Power Output (watts) |
| :---: | :---: | :---: | :---: |
| 15 | 0.125 | 157 | 0.186 * |
| 16 | 0.125 | 174 | 0.206 |
| 19 | 0.133 | 180 | 0.227 * |
| 20 | 0.133 | 179 | 0.225 |
| 22 | 0.148 | 197 | 0.276 |
| 23 | 0.157 | 187 | 0.278 * |
| 26 | 0.157 | 217 | 0.323 |
| 27 | 0.157 | 200 | 0.297 * |
| 29 | 0.167 | 202 | 0.319 * |
| 30 | 0.167 | 217 | 0.343 |
| 33 | 0.167 | 200 | 0.316 * |
| 34 | 0.175 | 219 | 0.363 |
| 36 | 0.184 | 226 | 0.394 |
| 37 | 0.192 | 218 | 0.396 * |
| 40 | 0.204 | 229 | 0.442 |
| 41 | 0.204 | 223 | 0.431 * |
| 43 | 0.214 | 226 | 0.468 * |
| 44 | 0.223 | 231 | 0.488 |

* housed in run wheel for that day.

Addendum

During the review of this thesis, it was suggested that the group regression lines for each experimental group be calculated in a different manner. Since the slopes of the individual regression lines (Appendix E) in each group are very similar, they were averaged and the group regression line formed. This method seperated the groups (Addendum Fig. 1) to a greater extent than the original procedure (Fig. 9). The standard error of estimate for each group was also reduced.

The conditioned-run wheel (Cl) group was significantly
different from all other groups. The conditioned (C2) and exercisedrun wheel (El) groups showed no significant difference. This suggests that the voluntary run wheels imposed a stress that produced a similar conditioning protocol used in the present study. This method also reinforces the possibility that at the higher power outputs, the conditioned rats are able to meet the increased demands through increased aerobic metabolism, whereas, the non-conditioned animals derived a greater percentage of the required energy to perform the work from anaerobic sources.

Addendurn


Addendum Summary of the test for homogenity of regression of sedentary control, exercised and conditioned groups

| Source of <br> Variation | DF | SS | MS | P |  |
| :--- | :---: | :---: | :---: | :---: | :---: |
| Groups | 4 | 0.0006544 | 0.0001636 | 2.70 | 0.05 |
| Error | 81 | 0.0007415 | 0.0000092 |  |  |

```
F
```

Newman-Keuls comparison between ordered means for regression slopes of sedentary conrol, exercised and conditioned rats

|  | SC | E2 | C2 | El | Cl |
| :---: | :---: | :---: | :---: | :---: | :---: |
| Means | 0.0165 | 0.0247 | 0.0285 | 0.0324 | 0.0386 |
| 0.0165 | 0.00 | 0.0082** | $0.0120 * *$ | 0.0159** | 0.0221** |
| 0.0247 |  | 0.00 | 0.0038 | $0.0077 * *$ | $0.0139 * *$ |
| 0.0285 |  |  | 0.00 | 0.0039 | $0.0101 * *$ |
| 0.0324 |  |  |  | 0.00 | $0.0062^{*}$ |
| 0.0386 |  |  |  |  | 0.00 |
| $\mathrm{R}=$ |  | 2 | 3 | 4 | 5 |

The Multiplier is 0.0014599
** $\mathrm{p}<0.01$

* $\mathrm{p}<0.05$

B30223


[^0]:    ? not reported

    * justification of conditioning program
    \# 12 intervals of running at $42 \mathrm{~m} / \mathrm{min}$., each lasting 30 sec., spaced 10 min. apart
    +2 min. rest periods every 10 min.

[^1]:    The relationship between heart rate and power output for the different
    experimental groups

[^2]:    $F_{0.01}(4,19) 4.50$

[^3]:    $F_{0.05}(4,15) 3.06$

