Calorimetric monitoring of the effect of endurance training and testosterone treatment on rat serum denaturation transition

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Abstract Differential scanning calorimetry (DSC) was applied to evaluate whether endurance training and/or testosterone treatment affect thermal behavior of rat serum. Thirty-six adolescent male Wistar rats were divided into six groups: three untrained and three trained groups, receiving weekly 0, 8, or 80 mg kg⁻¹_{BM} of testosterone enanthate (TE) each. Total testosterone was significantly higher in both groups treated with the higher TE dose. The rats' body mass and the levels of serum glucose, uric acid, and free fatty acids were additionally determined. The effect of TE treatment was clearly visible in DSC curves for sedentary rats and much less for trained rats. The differences between averaged endothermic denaturation transitions of serum from sedentary and physically active untreated groups were within an experimental error. A statistically significant increase of the width of peak at its half height and the ratio of excess heat capacity at 75 °C and in the temperature of peak maximum (C_{p75}/C_{pmax}) were observed due to TE treatment. The same two parameters were significantly lower for physically active rats. Significantly lower values of the enthalpy of serum denaturation (ΔH) and the first moment of the transition with respect to the temperature axis (M_1) were observed for endurance-trained than for sedentary rats.

Keywords Differential scanning calorimetry (DSC) · Protein denaturation · Rat serum · Testosterone treatment

Introduction

Serum is the specimen of choice for analysis in clinical chemistry. The proteome of serum is an indicator of the condition of the whole organism. Serum contains 60–80 mg of protein/mL and various small molecules including salts, lipids, amino acids, and sugars. The concentration range of plasma/serum proteins encompasses at least nine orders of magnitude [1]. The major protein constituents of serum include albumin (35–50 mg mL⁻¹), immunoglobulins, transferrin, haptoglobin, and lipoproteins. The thermally induced conformational transitions of proteins can be successfully investigated using differential scanning calorimetry (DSC). Recently, the DSC approach was applied to blood serum studies as a novel tool for disease diagnostics [2–9].

The possibility of DSC application in sport medicine has been also suggested and the effect of physical effort on the serum thermal profile of young male amateur cyclists has been investigated [10]. Here, we explore this approach for studying the effect of endurance training and testosterone treatment on rat serum. Calorimetry has already been applied in studies of newborn rat blood serum after maternal administration of cyclophosphamide, an anti-cancer chemotherapy drug [11]. However, it should be expected that “a thermodynamic signature” for the serum of adult rats differs from that of a newborn's because a sharp difference has been detected between protein patterns of rat serum samples of newborn and adult animals [12].

The trainability of individuals is related to the testosterone and hormonal parameters associated with this

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hormone, such as the testosterone/sex hormone-binding globulin (SHBG) ratio and the testosterone/cortisol ratio [13]. Exercise training is often combined with anabolic androgen use because the suitable level of circulating testosterone is important for an appropriate anabolic environment. About 60 % of circulating testosterone is strongly bound to SHBG, about 40 % of the hormone is bound to albumin, and only 1–4 % exists as free hormone. The biologically active testosterone includes both the free and albumin-bound fractions because the binding between testosterone and albumin is weak. Despite an increasing knowledge related to the acute and chronic testosterone responses to physical exercise and training, data on the changes in testosterone concentration after endurance and strength training still remain controversial.

Acute physical exercise, due to its increase in oxygen consumption, promotes the increase of the oxygen free radical (OFR) formation. When the OFR production exceeds the tissue's antioxidant ability, the result is oxidative stress. Physical training is able to induce adaptational changes in the body capable of preventing the harmful effects of OFR [14]. It was reported that uric acid (UA) reduces exercise-induced oxidative stress. High UA concentrations could confer protection against free radical activity in vivo [15]. Among other biochemical markers of exercise metabolic effects are glucose, lactate, triglyceride, glycerol, and fatty acids.

The aim of this study was to monitor changes in DSC denaturation profile of rat serum due to endurance training and testosterone treatment. In addition, the relationships between several biochemical markers of exercise metabolic stress and parameters of serum thermal transition were discussed.

Experimental

Groups of animals

Adolescent male Wistar rats of 90–100-g initial body mass (BM), from the stock of the Mossakowski Medical Research Center (Warsaw, Poland), were used for the study. The rats were housed four to five per cage, at a 12-h light/12-h dark cycle, 22–24 °C ambient temperature, and 45–65 % relative humidity, and were allowed ad libitum access to standard pellet rat chow and tap water.

Before the start of the study, all rats were run-tested on a motorized rodent treadmill (3 × 5 min, with 15-min breaks) to habituate them to the training and to eliminate rats unwilling to run. The selected rats were randomized between the following groups: untrained controls (UTr, $N = 6$), untrained rats receiving 8 mg kg⁻¹ BM (UTr + TE8, $N = 6$) or 80 mg kg⁻¹ BM (UTr + TE80, $N = 6$) of testosterone enanthate (TE) weekly, endurance-trained steroid-

untreated rats (Tr, $N = 6$), and endurance-trained rats receiving 8 mg kg⁻¹ BM (UTr + TE8, $N = 6$) or 80 mg kg⁻¹ BM (Tr + TE80, $N = 6$) of TE weekly.

Stock TE solution (*Testosteronum enanthatum*, Jelfa, Poland) was diluted with sesame oil as needed and injected intramuscularly once a week for 6 weeks, alternately into the right or left hindlimb. TE-untreated rats were given the same volume of the oil according to the same schedule.

The study protocol was approved by the First Warsaw Ethical Committee for Animal Experiments of the Polish Academy of Sciences, Warsaw, Poland (Certificate of approval No. 251).

Endurance training

The rats scheduled for endurance training were exercised on a rodent treadmill (at 0° inclination) 5 days a week, for 6 weeks. The treadmill speed was 16 m min⁻¹ during the first week, was increased by 4 m min⁻¹ weekly over the next 3 weeks, and was kept at 28 m min⁻¹ for the remaining training sessions. The session duration started at 40 min day⁻¹ and was increased by 5 min daily during the first 4 weeks; during the last 2 weeks, the rats ran for 60 min daily at 28 m min⁻¹ [16, 17]. Two days after completion of the training, the rats were euthanized by decapitation.

Serum samples

Trunk blood was collected, allowed to clot at room temperature, and spun to obtain serum for total testosterone (TT) assays. Serum TT was determined with a radioimmunoassay (KIP19000, DIASource Immunoassays S.A., Belgium). In addition, the concentrations of serum glucose, uric acid (UA), and free fat acids (FFA) were assessed by enzymatic methods using commercially available diagnostic kits (Randox UK, cat. no. GL 2623, UA 230, and FA 115, respectively). The mean values in each group are shown in Table 1.

DSC measurements

Differential scanning calorimetry experiments were carried out on the blood serum obtained from rats belonging to one of the six mentioned groups. Serum samples were stored at -20 °C before the analysis. For DSC measurements, the samples were diluted 20-fold in degassed DDW. The pH of the diluted samples was within the range of 6.5–7.5.

DSC scans were performed using the VP DSC ultra-sensitive microcalorimeter (MicroCal Inc., Northampton, MA) with cell volume of 0.5 mL. The samples were heated from 20 °C to 100 °C with a scan rate of 1.0 °C min⁻¹. In addition, a constant pressure of about 1.8 atm was exerted on the liquids in the cells. The calorimetric data were

Table 1 Effects of testosterone enanthate (TE) treatment and endurance training on body mass (BM), serum total testosterone (TT) glucose, uric acid (UA), and free fatty acids (FFA) levels in male Wistar rats

Variable	UTr	UTr + TE8	UTr + TE80	Tr	Tr + TE8	Tr + TE80
BM/g	328.2 ± 29.8	319.0 ± 44.6	264.5* ± 8.3	297.2 ± 64.1	297.3 ± 33.1	264.8 ± 18.8
TT/nmol L ⁻¹	2.85 ± 2.76	2.69 ± 1.09	30.22*** ± 9.24	3.44 ± 3.14	2.88 ± 1.76	32.18*** ± 2.59
Glucose/mg dL ⁻¹	139 ^{###} ± 13.4	114.7 [#] ± 8.1	89.8 ± 6.9	106.9 ± 19.8	101.7 ± 9.2	95.3 ± 9.5
UA/mg dL ⁻¹	2.3 ± 0.4	2.1 ± 0.5	2.4 ± 0.5	2.3 ± 0.5	3.3 ⁺ ± 0.8	2.8 ± 0.7
FFA/mmol L ⁻¹	0.57 ± 0.16	0.68 ± 0.15	0.67 ± 0.26	0.63 ± 0.31	0.55 ± 0.19	0.83 ± 0.24

Data are means (±SD). UTr untrained rats, Tr endurance-trained rats, TE 8 mg kg⁻¹ BM TE once a week, TE80 80 mg kg⁻¹ BM TE once a week; * $p < 0.05$ versus the UTr and UTr + TE8 group, ** $p = 0.01$ versus Tr + TE8 group, *** $p < 0.001$ versus the respective untreated and TE 8 mg kg⁻¹ BM group, # $p < 0.05$, ### $p < 0.001$ versus UTr + TE80 group, + $p < 0.05$ versus untreated group (Kruskal–Wallis test)

corrected for the instrumental baseline water–water. DSC curves were normalized for the gram mass of protein and next a linear baseline was subtracted. The curves of apparent excess heat capacity, C_p^{ex} , versus temperature were plotted. Protein concentrations were determined according to its amount in dry mass of the serum sample.

Analysis

DSC curves were analyzed with the use of MicroCal Origin software. The following thermodynamic parameters have been found: the temperature of peak maximum (T_m), the enthalpy of serum denaturation (ΔH) (calculated as the area under the peak), the width of peak in its half height (HHW), and the normalized first moment (M_1) of the thermal transition with respect to the temperature axis calculated as

$$M_1 = \frac{\int_{T_i}^{T_f} T C_p dT}{\int_{T_i}^{T_f} C_p dT} \quad (1)$$

where T_i and T_f are the beginning and the end of denaturation, respectively. In addition, the ratios $C_{p75}/C_{p\text{max}}$ (where C_{p75} is the value of C_p^{ex} at 75 °C (the approximate temperature of the shoulder) and $C_{p\text{max}}$ the value of C_p^{ex} at T_m) were calculated.

Statistical analysis of the results was done with Statistica 10 using ANOVA, the U Mann–Whitney, or Kruskal–Wallis tests. The Shapiro–Wilk and the Levene tests were used to check the normality of the distributions and homogeneity of the variance, respectively. The significance of between-group differences was tested using a 2-way ANOVA, with training status and TE dose as the main factors, followed by the post hoc Tukey's HSD (honestly significant difference) test. In all cases, $p < 0.05$ was considered significant.

Results and discussion

The averaged DSC serum curve for the untrained control group of rats is shown in Fig. 1. The shaded area represents

Fig. 1 Mean serum DSC curve ($n = 6$) of rats from untrained control group. The shaded area is the standard error of the mean at each temperature

the standard error of the mean at each temperature. The complex endothermic transition observed in the temperature range 40–90 °C arises from the unfolding of proteins within the serum. Each individual protein has a unique thermal profile under certain experimental conditions. The peak representing serum denaturation is formed as the weighted sum of thermal profiles of constituent proteins, but the overall shape of the denaturation peak may be modified by binding interactions and environmental factors [18].

In general, there are some similarities between denaturation DSC profiles of rat serum and human plasma/serum of healthy individuals [2–6]. The temperature range of the whole transition is practically the same; however, the endothermic peak is more symmetric for rat than for human serum. The rat DSC profile (Fig. 1) is surprisingly simple and component transitions are not well resolved. The major transition represented by a clear peak with T_m ca. 66 °C is not as sharp as that observed at ca. 62 °C for human serum [2–7]. Two shoulders at about 75 °C and 82–83 °C are barely visible in Fig. 1. It is not clear which rat serum proteins contribute to the major transition and shoulders.

Fig. 2 The comparison of averaged DSC curves of serum for rats from untrained (*solid line*) and endurance-trained (*dot line*) steroid-untreated groups

DSC curves of individual proteins in human plasma, useful in the analysis of plasma or serum transition, have been presented by Garbett et al. [2], but for rat proteins there is no corresponding set of data. Notwithstanding this, some analogies seem to be possible.

The main protein fractions of human as well as rat blood serum are albumins and globulins. A detailed investigation of rat serum proteome by Gianazza et al. [12] shows, however, that the protein assortment in rat serum is definitely different from that in human serum. Some proteins are present in much lower concentration in rat than in human serum (e.g., orosomucoid or α_1 -acidic glycoprotein and haptoglobin under physiologic conditions), while others are present in much higher concentration (e.g., C-reactive protein); some proteins are species specific to the rat. Even common proteins display different physico-chemical parameters [12]. The denaturation behavior of albumin, the most abundant serum protein, depends on the species. Significant differences have been reported with respect to the conformational stability and the mechanism of the unfolding pathway for mammalian albumin [19, 20]. The calorimetric enthalpy as well as thermal denaturation temperature has been shown lower for rat than for human and bovine defatted albumin [19]. It should also be noted that the transition temperatures for fatty acids free albumins are lower than for their non-defatted counterparts [20].

Fig. 2 compares the averaged DSC curves of blood serum for rats from the untrained and endurance-trained steroid-untreated groups. These mean curves are very similar. The slight decrease of transition peak intensity seems to be within the experimental error. However, a distinct feature of serum transition observed for the Tr group is the lack of the shoulder at 75 °C.

The effect of testosterone treatment on the serum denaturation transition is shown in Fig. 3 and 4 for the untrained and endurance-trained group, respectively. A

Fig. 3 The averaged serum DSC curves for rats from UTr, UTr + TE8, and UTr + TE80 groups

Fig. 4 The averaged serum DSC curves for rats from Tr, Tr + TE8, and Tr + TE80 groups

decrease of intensity and a broadening of the thermal transition peak with increasing of testosterone dose can be seen. It is interesting that these effects are more clearly visible in the group of untrained rats. Additionally, for TE-treated sedentary rats, the shoulder at about 75 °C is well visible (Fig. 3). Surprisingly, the intensity of the averaged DSC curve for the serum from trained rats treated with the higher testosterone dose (Tr + TE80) seems to be more similar to the intensity of the thermal curve received for the steroid-untreated group (Tr) than for that treated with a lower TE dose (Tr + TE8). For the Tr + TE80, but not for the Tr + TE8 group, the shoulder in the 75–80 °C temperature range occurs (Fig. 4).

Such results suggest that the effects of testosterone treatment and physical activity may be antagonistic to each other. Recently, similar antagonistic effects of testosterone and physical activity on muscle fiber diameter in rats were reported [21].

The comparison of thermodynamic parameters summarized in Table 2 suggests that there are no significant

Table 2 Effects of testosterone enanthate (TE) treatment and endurance training on thermodynamic parameters of serum denaturation transition

Variable	UTr	UTr + TE8	UTr + TE80	Tr	Tr + TE8	Tr + TE80
$T_m/^\circ\text{C}$	66.2 ± 0.6	65.8 ± 0.4	65.4 ± 1.0	66.4 ± 1.3	65.5 ± 0.6	65.9 ± 0.4
$\Delta H/\text{J g}^{-1}$	18.8 ± 3.4	18.8 ± 2.5	16.9 ± 1.8	16.7 ± 1.9	15.2 ± 1.8	16.5 ± 1.4
HHW/ $^\circ\text{C}$	$20.1^{**} \pm 1.5$	23.4 ± 3.2	26.1 ± 1.8	19.5 ± 3.1	22.1 ± 3.1	23.4 ± 2.0
$M_1/^\circ\text{C}$	68.0 ± 0.7	68.1 ± 0.6	68.8 ± 0.8	67.7 ± 0.9	67.5 ± 0.6	67.9 ± 0.6
C_{p75}/C_{pmax}	$0.62^{**} \pm 0.04$	$0.75^* \pm 0.12$	0.80 ± 0.08	$0.57^{**} \pm 0.08$	0.64 ± 0.07	0.75 ± 0.09

Data are means ($\pm$ SD). UTr untrained rats, Tr endurance-trained rats, TE 8 mg $\text{kg}_{\text{BM}}^{-1}$ TE once a week, TE80 80 mg $\text{kg}_{\text{BM}}^{-1}$ TE once a week; * $p < 0.05$ and ** $p < 0.005$ versus the respective TE 80 mg $\text{kg}_{\text{BM}}^{-1}$ group (post hoc Tukey test)

differences between the transition temperatures T_m for various subgroups. The results presented in this table indicate that the HHW and C_{p75}/C_{pmax} ratio have been increased by TE treatment and decreased by endurance training. Two-way ANOVA has revealed that these effects are statistically significant (for both parameters, $p < 0.001$ in the case of TE treatment factor and $p < 0.05$, $p < 0.01$ for HHW and C_{p75}/C_{pmax} values, respectively, in the case of endurance training factor). No significant interactions between these factor effects have been found. The Tukey post hoc test for C_{p75}/C_{pmax} ratio has yielded significant differences between TE-untreated rats and those receiving the higher TE dose both in the sedentary and trained groups. For the HHW parameter, essential difference has been obtained between the UTr and UTr + TE80 groups. Additionally, the results of statistical analysis indicated significantly lower values of ΔH as well as M_1 for the endurance-trained in comparison with sedentary rats ($p = 0.004$ and $p = 0.009$, respectively). No significant effect of TE dose on these values was observed.

Statistical correlations found between some biochemical and thermodynamic parameters, presented in Tables 1 and 2, may be helpful in the explanation of DSC results. The positive moderate correlation between FFA level and C_{p75}/C_{pmax} ratio ($r = 0.48$, $p = 0.001$) suggests a relationship between the appearance of the shoulder at about 75°C and the level of fatty acids in serum. A migration of fatty acids, their release or binding to albumin, changes the protein thermal profile of denaturation. As albumin is the most abundant serum protein, the DSC profile of serum should depend highly on denaturation transition of albumin, and hence on the ratio of fatty acid free albumin and fatty acid-carrying albumin fractions. The DSC denaturation transition for the albumin with bound fatty acids is strongly shifted to the higher temperature in relation to the defatted albumin [20, 22]. Thus, a decrease of fatty acid free (defatted) fraction of albumin should lead to a reduction of the main peak intensity. On the other hand, an increase in the number of albumin molecules with bound fatty acids (fatty acid carrying fraction) should cause an increase of

peak intensity in the higher temperature range, above the transition temperature. Therefore, the total effect can be seen as a broadening of DSC serum denaturation peak and an increase of HHW and C_{p75}/C_{pmax} values. Just such an effect has been observed with the TE dose increase in our study. The results shown in Table 1 indicate a significantly higher TT level for both TE 80 mg $\text{kg}_{\text{BM}}^{-1}$ groups—sedentary and endurance-trained. However, the FFA level was slightly higher only in the Tr + TE80 group.

It is well established that free fatty acids in blood serum are mainly derived from the breakdown of body fat stores, which can be mobilized by catecholamines and exercise. Hydrolysis of the intracellular TG is regulated by hormone-sensitive lipase (HSL), the enzyme expressed in adipose tissue, in cardiac and skeletal muscle cells, preferentially in oxidative fibers [23]. Langfort et al. [24] have reported that testosterone may increase HSL activity, thus mobilizing cellular TG stores, while carbohydrate oxidation is diminished. Circulating FFAs are known to reach a maximum shortly after the cessation of exercise, which is a consequence of their augmented release into the plasma pool during exercise, subsequently followed by augmented removal of FFA during the recovery [25]. This may most likely explain why the FFA level in serum samples collected after complete recovery from trained rats given the lower TE dose was moderately lower than in any other group of animals. Unfortunately, any significant differences have not been found between FFA levels in the various rat groups (Table 1). A slightly suppressed level of TT in the group of trained rats treated with a lower TE dose (8 mg $\text{kg}_{\text{BM}}^{-1}$) as compared to trained, steroid-untreated controls might result from exogenous administration of TE, which is known to induce suppression of endogenous testosterone [26, 27]. Notably, for this group, a disappearance of a shoulder at about 75°C on the DSC curve was observed.

The relationship between FFA and TT levels seems to be complicated and changed by a physical activity. In our experiment, a moderate positive correlation ($r = 0.49$, $p = 0.02$) was found between FFA and TT values for

endurance-trained rats, but not for untrained animals. Likewise, the statistically significant, positive correlations between FFA level and such thermodynamic parameters as ΔH ($r = 0.50$, $p = 0.02$), HHW ($r = 0.49$, $p = 0.02$), and M_1 ($r = 0.49$, $p = 0.02$) were recorded for the trained, but not for the untrained groups. No significant correlation between the above-mentioned parameters and TT levels in trained rats were found.

It is interesting that body mass BM was negatively and TT level was positively correlated with HHW ($r = -0.47$, $p = 0.003$ and $r = 0.48$, $p = 0.002$, respectively). However, it is not clear which of these correlations can indicate potential causal relationships. There is an important relationship between BM and serum TT level. The significant negative correlation between these variables shown in a previous investigation [28] has been confirmed by the results of our current study ($r = -0.52$, $p < 0.001$).

The glucose level (see Table 1) has been markedly decreased by both studied factors: endurance training ($p < 0.005$) and by TE treatment ($p < 0.001$). However, for TE treatment effect, the differences between the trained groups have been found not statistically significant. In addition, the mean glucose level for rats receiving $80 \text{ mg kg}_{BM}^{-1}$ was very similar in the sedentary and trained groups. A negative correlation ($r = -0.64$, $p = 0.001$) between the glucose level and C_{p75}/C_{pmax} ratio occurs for untrained and also for trained rats. For untrained, but not for trained rats, significant negative correlations have been obtained between the glucose level and HHW ($r = -0.76$, $p < 0.001$), as well as between the glucose level and M_1 ($r = -0.42$, $p = 0.046$).

High UA concentrations are associated with increased serum antioxidant capacity and reduced oxidative stress during acute physical exercise in healthy subjects [15]. In our experiment, the level of uric acid (UA) varied slightly among different groups of rats (see Table 1). There was no difference among the untreated groups of rats (UTr and Tr). However, the U Mann–Whitney test revealed significantly higher UA level for the trained, compared to the untrained rats ($p = 0.01$). It is in agreement with the finding that the extended acute exercise is associated with a temporary increase of uric acid produced when a relatively high exercise intensity and a high rate of energy consumption trigger activation of adenylate kinase and AMP signaling networks to insure cellular energy homeostasis [14, 29, 30]. The highest value of UA was recorded for the Tr + TE8 group. The significant difference between the Tr + TE8 and Tr groups was found ($p = 0.01$) and no effect of TE treatment on UA level was observed in the sedentary group. It is intriguing that a relatively high positive correlation was found between UA level and M_1 ($r = 0.71$, $p = 0.001$) for untrained, but not for trained rats.

Conclusions

The effects of chronic treatment with supraphysiological TE doses, given alone or in combination with moderate intensity endurance training, on rat serum denaturation transition were studied using DSC. Significantly higher serum total testosterone levels were recorded only in both trained and untrained groups given the higher TE dose. A decrease of intensity and a broadening of the thermal transition peak with increasing doses of exogenous testosterone were observed in the untrained groups of animals. For trained rats, the analogous tendency was not so clear, probably due to some kind of antagonistic effects of testosterone and exercise training on serum composition, protein conformational changes, or protein–protein and protein–ligand interactions within the rat serum. Opposite effects of testosterone treatment and training on some transition parameters, such as HHW and C_{p75}/C_{pmax} ratio, were observed. Endurance training was associated with statistically significant decrease, while TE treatment with an increase in HHW and C_{p75}/C_{pmax} ratio. The results of statistical analysis performed for all groups showed that values of the enthalpy of transition, ΔH , and the first moment temperature, M_1 , were also lower for the trained in comparison with untrained rats. The first moment temperature was reported as a very useful parameter in disease diagnosis based on serum DSC studies [2–6]. Lower M_1 values for healthy individuals and an increase of M_1 with the severity of the disease were observed. Lower M_1 values for trained rats suggest their better health status compared to their sedentary counterparts.

A significant relationship was found between the level of free fatty acids in serum and an increase in the intensity of the shoulder at about 75°C (positive correlation between FFA and C_{p75}/C_{pmax} ratio). Moderate positive correlations between FFA level and such thermodynamic parameters as ΔH , HHW, and M_1 were shown for the trained, but not for sedentary (untrained) rats. Just for endurance-trained rats, moderate positive correlation between serum FFA and TT levels was present. Glucose level, decreased by endurance training and due to TE administration, was negatively correlated with C_{p75}/C_{pmax} ratio. For sedentary rats, negative correlations were observed between the glucose level and HHW or M_1 and relatively high positive correlation between UA level and M_1 . The interpretation of these interesting results is not entirely clear-cut at present and remains to be validated with a larger group of rats in future investigations.

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