Maintenance on a ketogenic diet: voluntary exercise, adiposity, and neuroendocrine effects

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Abstract

Introduction—Adherence to low-carbohydrate, ketogenic diets has been associated with greater weight loss in the short-term than low-fat, calorie-restricted diets. However, consumption of ketogenic diets may result in decreased voluntary exercise and thus render long-term weight loss and maintenance of weight loss difficult.

Methods—Rats were maintained on either a non-ketogenic chow (CH) diet or a low-carbohydrate, ketogenic diet (KD) for 6 weeks. Half of each dietary group was sedentary, while the other half was given access to a running wheel. Running wheel activity (total distance and meters/minute), plasma leptin and insulin, adiposity, and hypothalamic mRNA for neuropeptide Y (NPY) and proopiomelanocortin (POMC) were measured to assess activity-related effects in animals maintained on KD.

Results—With access to a running wheel, rats on KD engaged in similar levels of voluntary activity as CH rats and both dietary groups decreased caloric intake. Caloric intake increased over time such that it was significantly greater than sedentary controls after one month of access to the wheels, however body weight remained decreased. Sedentary rats maintained on KD had increased adiposity and plasma leptin levels and decreased hypothalamic POMC mRNA, as compared to sedentary CH rats. KD rats with access to a running wheel had similar levels of adiposity and plasma leptin levels as CH rats with access to running wheels, but significantly increased POMC mRNA in the arcuate.

Conclusion—We demonstrate that maintenance on KD does not inhibit voluntary activity in a running wheel. Furthermore, prevention of KD-related increased adiposity and plasma leptin, as measured in sedentary KD rats, significantly increases levels of the anorexigenic neuropeptide POMC mRNA.

Keywords
proopiomelanocortin; leptin; low-carbohydrate diet; ketosis; running wheel
Introduction

The prevalence of overweight and obesity among adults in the United States has increased significantly over the past four decades (1). Numerous diets have been proposed to induce weight loss, and it is estimated that between 30 and 45% of adults in the United States use specific dietary approaches in an attempt to lose weight (2). Although low fat, reduced-calorie diets coupled with increased energy expenditure are generally recommended for weight loss, the use of low-carbohydrate diets has increased in recent years (3). Proponents of low-carbohydrate, ketogenic diets promote the concept that caloric intake and dietary fat are not the culprits in obesity, and that ultimately minimization of carbohydrate intake leads to weight loss, decreased hunger, and maintenance of reduced body weight (4, 5). Low carbohydrate diets have, in fact, been associated with greater weight loss in the short-term as compared to weight loss associated with low fat diet users (6–11). However, long-term evaluations found no significant differences in weight loss between low-carbohydrate dieters and controls at one year (11, 12).

While many individuals engage in dieting strategies in an attempt to lose weight, increased physical activity is an important component of the weight-loss equation. Studies comparing weight loss with diet alone as compared to those that included increased physical activity demonstrate increased weight-loss maintenance in individuals who regularly exercised (13, 14). Individuals consuming carbohydrate-restricted diets report fatigue and reduced vigor in response to exercise significantly more than individuals consuming high-carbohydrate diets (11, 15). Furthermore, maintenance on a ketogenic diet results in reduced free-living physical activity, a response that can be counter-productive in individuals trying to lose weight (16). Fatigue and perceived effort during exercise have been compared in trained, overweight adults adhering to a low-carbohydrate diet that was ketogenic or to a control diet that was also low in carbohydrate but non-ketogenic (5% vs. 40% carbohydrate). While weight loss was similar among diet groups, there was a positive correlation between feelings of fatigue while exercising and blood ketone concentrations. These data suggest that ketogenic diets enhance fatigability and can reduce the desire to exercise (16).

Given reported differences in levels of physical activity by individuals consuming low carbohydrate diets (16), the increased fat content of low carbohydrate diets (8, 17) and the known effects of consuming a high fat diet on adiposity (18–23), we hypothesized that rats maintained on a ketogenic diet would engage in less voluntary activity when given access to a running wheel than chow-fed rats. We evaluated running wheel activity, caloric intake and body weight in rats given access to a running wheel for four consecutive weeks, as well as adiposity and plasma insulin and leptin. We have previously reported that maintenance on a ketogenic diet affects neuropeptide Y (NPY) and proopiomelanocortin (POMC) mRNA in the arcuate nucleus of the hypothalamus (24), therefore we included measures of these parameters after exposure to running wheels.

Methods

Thirty-two male Long Evans rats (Harlan, Indianapolis, IN) weighing 225–250 g were individually housed in clear plexiglass tub cages and maintained at a constant temperature.
Food intake, body weight, and running wheel access

After one week of acclimation to the laboratory, during which time rats were given ad libitum access to rodent chow (Harlan Teklad 2018 18% protein diet, 3.4 kcal/g), rats were weight-matched and divided into two groups. The first group maintained ad libitum access to chow (CH) and the second group was placed on a ketogenic diet (KD) from which 5% of the calories were derived from carbohydrate, 80% from fat, and 15% from protein (Table 1, Research Diets D06040601, 6.1 kcal/g, Research Diets Inc., New Brunswick, NJ). Caloric intake and body weights were measured and recorded daily.

Following one week of access to CH or KD, rats were weight-matched within their diet groups. Running wheels were unlocked for half of each group at this point (CH-RUN (n=8) and KD-RUN (n=8)), and remained locked for the other half of the rats who served as sedentary controls for each diet group (CH-SED (n=8) and KD-SED (n=8)). For rats with running wheel access, running wheel activity was computer monitored consecutively and recorded every 30 minutes for 30 days.

Blood and Tissue Collection

At the end of the experiment, rats were killed between 12:00 and 2:00 PM by decapitation under ether inhalation anesthesia. Brains were rapidly removed, submerged into iced isopentane for 25 seconds and immediately stored in dry ice. Fifty μl of trunk blood was collected for β-hydroxybutyrate analysis (Stat-site ketosis meter, GDS Diagnostics, Elkhart, IN), and the remaining trunk blood was collected into chilled K+EDTA vacutainer tubes, briefly placed in ice, and then centrifuged at 4°C for 15 m at 2000 rpm. Plasma was aspirated into eppendorf tubes. Blood and brains were stored at −80°C until processing. Epidydimal, subcutaneous, and retroperitoneal fat pads were also removed and weighed.

Radioimmunoassay

Radioimmunoassays (Millipore, St. Charles, MO) were used to determine baseline levels of plasma insulin and leptin. The rat insulin RIA kit had a sensitivity of 0.1 ng/ml. The intra- and interassay coefficients of variation (CV) were below 2.2% and 9.4%, respectively, and the correlation coefficient for the standard curve was 0.85. The rat leptin RIA sensitivity was 0.5 ng/ml. The intra- and interassay CV were below 2.4% and 5.7%, respectively, and the correlation coefficient for the standard curve was 0.94. Volumes of 100 μl of plasma were used in duplicate samples for each assay, as directed by the manufacturer.

In situ hybridization

Brains were coronally sectioned at 14 μm, mounted onto electrostatically charged Superfrost Plus slides (Fisher Scientific), and stored at −80°C. Brain slices were fixed with 4%
paraformaldehyde and dehydrated with an ascending series of alcohols. Sections from each rat containing the arcuate nucleus of the hypothalamus were selected and stored at −80°C for future processing. Plasmids of NPY and POMC were linearized with the appropriate restriction enzymes. Antisense riboprobes were labeled with 35S-labeled UTP (Perkin Elmer), using in vitro transcription systems with T7 polymerase, according to protocols provided by the manufacturer (Promega). Probes were then purified using Quick Spin RNA columns (Roche Diagnostics).

For processing, slides were warmed and rinsed in triethylamine (TEA) buffer (pH 8.0) and TEA with acetic anhydride. Sections were incubated in hybridization buffer comprised of 50% formamide, 0.3 M NaCl, 10 mM Tris HCl (pH 8.0), 1 mM EDTA (pH 8.0), 1 × Denhardt’s solution (Eppendorf), 10% dextran sulfate, 10 mM DTT, 500 μg/mL yeast tRNA, and 10^8 cpm/μL 35S-UTP, and incubated overnight in a 56°C humid chamber. After hybridization, sections were washed three times in 2X SSC followed by one wash in 2X SSC + DTT at 56°C. Slides were then treated with 20 μg/mL RNase A (Sigma) in buffer containing 5 M NaCl, 0.5 M EDTA, 1 M Tris, pH 7.5 and ddH2O. Sections were washed twice in 2X SSC + DTT, and then twice in 0.1X SSC + DTT, and dehydrated in an ascending series of alcohols. Slides were exposed to Kodak Biomax film for 2 days. Autoradiographic images were then scanned, and quantified with Scion Image software (National Institute of Health), utilizing autoradiographic 14C-microscales (Amersham Pharmacia Biotech) as a standard. An experimentally blinded observer performed scanning and quantification. Data for each animal were means of the product of hybridization area × density, with the background density subtracted from the three sections, reflecting the region-specific levels of gene expression. Data for each animal were normalized to CH-SED controls as 100%, and are expressed as mean ± standard error (SEM).

Data analyses

Data are represented as mean ± SEM. Caloric intake and body weight were first analyzed by 2-way ANOVA for repeated measures (diet × exercise over time), followed by planned t tests. Activity data were analyzed by repeated measures ANOVA, with time as the repeated measure, and further analyzed with post hoc Bonferroni comparisons when significant differences were found. Because terminal body weights were different between the sedentary and running groups, fat pads were analyzed as percentage of overall body weight for each depot measured. These values, and the RIA data were assessed by ANOVA and further analyzed by Tukey’s Multiple Comparison Test where appropriate. A value of P < 0.05 was considered significant.

Results

Caloric intake, body weight and activity levels

Prior to gaining access to the running wheels, there were no differences in daily mean caloric intake between rats maintained on CH and those maintained on KD (79.7 ± 1.2 and 80.2 ± 1.9 kcals/day, respectively). As depicted in Figure 1, access to a running wheel resulted in decreased caloric intake in CH-RUN as compared to CH-SED in the first week (CH-SED: 84.7 ±1.6 and CH-RUN: 77.6 ± 1.2 kcals/day (P <0.01)). Mean daily caloric
intake during the second and third weeks of running wheel access did not differ between
CH-RUN and CH-SED rats, however at 4 weeks CH-RUN rats consumed significantly more
calories per day than did CH-SED rats (CH-SED: 86.2 ±1.7 and CH-RUN: 95.1 ±1.1
kcals/day (P <0.05)). Access to running wheels had a similar effect in KD-RUN, as
compared to KD-SED. After one week of access to the wheels, KD-RUN rats consumed
significantly fewer calories per day than KD-SED rats (KD-SED: 82.1 ±1.8 and KD-RUN:
73.6 ±2.3 kcals/day (P < 0.01)). This difference was abolished in the second week of
running and KD-RUN rats consumed significantly more calories per day than KD-SED rats
in the 4th week of access to the running wheels (KD-SED: 84.4 ± 2.0 and KD-RUN (P <
0.01)). There were no differences in mean daily caloric intake between CH-SED and KD-
SED or CH-RUN and KD-RUN at any of the weekly intervals. Cumulative caloric intake is
represented in Table 2. While differences in caloric intake were measured on a week-by-
week basis with access to a running wheel, cumulatively there were no differences between
groups.

Access to running wheels resulted in decreased body weight gain in CH-RUN and KD-RUN
rats, as compared to rats without access to wheels (Figure 2). After two weeks of access to
the running wheels, CH-RUN rats had gained significantly less body weight than CH-SED
rats (P < 0.05). This difference persisted for the duration of the experiment such that CH-
RUN body weights were 24.2% less than those of CH-SED. Similar effects were measured
in KD rats. Body weight gain in KD-RUN was significantly less than in KD-SED after two
weeks of access to the wheels, (P <0.01), and this difference persisted throughout the
experiment such that after one month, KD-RUN body weights were 30.8% less than KD-
SED. The type of diet on which the rats were maintained had no effect on body weight
change over time.

Activity levels were recorded every 30 minutes for the month during which rats had access
to the running wheels. As depicted in Figures 3A and 3B, CH-RUN and KD-RUN rats had
similar levels of activity (total distance (m)) and ran at similar rates (m/min) for the duration
of the experiment.

**Plasma profiles and body composition**

Maintenance on KD resulted in significantly increased levels of s-hydroxybutyrate, as
compared to rats maintained on CH. β-hydroxybutyrate levels were elevated in both KD-
SED (0.36 ± 0.07 mM) and KD-RUN (0.31 ± 0.04 mM), as compared to chow-fed rats (CH-
SED: 0.18 ± 0.02 and CH-RUN: 0.16 ± 0.02 mM (P <0.01 for both groups, data not
shown)). Rats maintained on KD (SED and RUN) had lower levels of plasma insulin than
chow-fed rats (P < 0.05 in both cases, Figure 4A). There were no running wheel-related
effects on insulin levels for either dietary group. Figure 4B demonstrates differences in
plasma leptin levels. Access to the running wheels resulted in decreased plasma leptin in
both CH-RUN and KD-RUN rats, as compared to their sedentary counterparts (P <0.05 in
both cases). After 1 month of access to the running wheels, CH-RUN leptin levels were
36.6% less than CH-SED. The difference was much greater in KD rats. Plasma leptin was
60.8% lower in KD-RUN than in KD-SED. While the KD-SED rats had significantly higher
levels of leptin than CH-SED, there were no diet-related differences after access to the
wheels such that CH-RUN and KD-RUN had similar plasma leptin levels. In sedentary rats, maintenance on KD resulted in increased epididymal fat, retroperitoneal and subcutaneous fat, as compared to maintenance on chow (Table 2, $P < 0.05$ in all cases). After 1 month of access to running wheels, CH-RUN and KD-RUN groups had levels of epididymal and retroperitoneal fat that were statistically indistinguishable from CH-SED controls, whereas subcutaneous fat was significantly less in CH-RUN rats ($P < 0.05$). Overall, KD-RUN rats had 47.7% less body fat than KD-SED ($P < 0.05$) after 1 month of access to the running wheels, and CH-RUN had 27.2% less total body fat than CH-SED.

**Hypothalamic gene expression**

In the arcuate nucleus of the hypothalamus, there were no differences in expression levels of mRNA for NPY between any of the dietary or exercise groups (Figure 5A). As for POMC, there was no effect of exercise in the CH-fed animals such that CH-SED and CH-RUN were not different from one another. Maintenance on KD, however, did affect levels of POMC mRNA expression. POMC mRNA expression was significantly lower in KD-SED, as compared to CH-SED ($P <0.05$), and was significantly greater in KD-RUN ($P <0.05$, Figure 5B).

**Discussion**

Studies in human subjects maintained on low carbohydrate diets suggest an association between these types of diets and a reduction in voluntary physical activity. In the current studies, we demonstrate that maintenance on a ketogenic diet does not result in decreased voluntary activity when rats were given access to a running wheel, as compared to activity levels of rats maintained on a non-ketogenic, chow diet. Running wheel access resulted in similar effects on body weight, caloric intake and activity levels in rats maintained on CH or KD. Interestingly, exercise in rats maintained on KD had a significant effect on hypothalamic POMC mRNA expression levels that was not observed in exercising rats maintained on chow.

In the sedentary groups, rats on KD consumed the same number of calories per day as CH controls. This finding is consistent with our previous work, in which rats maintained on a ketogenic diet had similar rates of weight gain and caloric intake as did chow-fed rats (24). Previous research has demonstrated that rats maintained on CH and given voluntary access to running wheels decrease food intake and body weight gain as compared to sedentary controls over a 7- to 10-day period (25–27) and that over a 4 week period, rats with access to a running wheel gain significantly less than sedentary rats (28). Whereas low-carbohydrate and ketogenic diets have been associated with increased fatigue in response to exercise and reductions in free- living physical activity in human subjects (16, 29, 30), our current results demonstrate that access to a running wheel induced changes in caloric intake and body weight, regardless of diet-type. After one week of access to a running wheel, both CH and KD rats had a significant decrease in daily caloric intake. This deficit was attenuated during the second week, and rats in both groups consumed more calories per day than did sedentary controls by the 4th week. Differences in body weight were evident after two weeks of wheel running. Despite increased caloric intake in the 4th week, body weight remained
significantly lower in CH-and KD-fed rats with access to wheels, as compared to their sedentary counterparts. This is likely due to the similarities in cumulative caloric intake over the course of the experiment, which demonstrates that overall, the reduced body weight in CH-RUN and KD-RUN is associated with increased physical activity.

Several lines of evidence indicate that chronic consumption of diets high in fat contributes to obesity. Body fat storage occurs at a greater rate when excess energy comes from fat than when it comes from carbohydrate or protein (19, 20). Animals maintained on high fat diets increase energy intake, increase efficiency of body fat gain, and become obese (23). Cross-sectional studies in humans have demonstrated that the concentration of dietary fat is correlated with increased body weight and body mass index (18, 21, 22). We have previously reported that rats on KD have increased adiposity despite no difference in body weight as compared to CH-fed rats (24). With the exception of subcutaneous fat, chow-and KD rats with access to running wheels had similar levels of body fat, as compared to sedentary, chow-fed rats. These data demonstrate that body fat deposition is reduced to the same degree in chow and KD, regardless of the increased dietary fat and propensity for fat storage in KD rats.

We have previously reported ketogenic diet-related effects on the central melanocortin system, including decreased POMC mRNA expression, increased sensitivity to the anorexigenic effects of agonism of the MC3/4R by central administration of MTII, and increased expression levels of the MC4R (24). In the current experiments, we replicate and extend findings on the effects of KD on the hypothalamic melanocortin system. In sedentary rats fed KD, there was less POMC mRNA expression, however after 4 weeks of voluntary running POMC was increased, as compared to sedentary chow and sedentary KD rats. Overall, there are numerous factors that influence central neuropeptide systems involved in the regulation of energy that have not been investigated in animals maintained on a ketogenic diet. Our current experiments allow for speculation about how exercise-induced changes in leptin and caloric intake may affect hypothalamic POMC mRNA expression levels in rats maintained on KD.

It is known that leptin receptors are expressed on POMC neurons, and leptin administration stimulates POMC mRNA expression in CH-fed animals (31, 32). Additionally, increased POMC expression has been demonstrated in cases of overfeeding and decreased voluntary caloric intake (33). This relationship is inconsistent with both sedentary KD, in which leptin levels are high and POMC expression is reduced, and active KD rats in which leptin levels are reduced and POMC expression is elevated. These data suggest the possibility of a diet-related disconnect in the relationship between peripheral leptin and the central melanocortin system, as it has been characterized in animals maintained on non-ketogenic diets. Whether or not this is the case is currently unknown and admittedly speculative, and POMC responsivity to circulating leptin in animals maintained on a ketogenic diet requires further investigation.

A second possible explanation for increased POMC expression in KD-RUN is that these rats consumed more kcals per day than did sedentary controls on the same diet. Increased POMC mRNA expression may be a reflection of this increased caloric intake, as rats that are
overfed display increased POMC mRNA expression (33). However, the pattern of intake, in terms of total kcals per day, was the same in CH-RUN and KD-RUN, yet there was no increase in POMC mRNA in CH-RUN. It is possible that this is due to differential sensitivity to increased caloric intake in rats consuming KD, as compared to those consuming CH, at the level of arcuate POMC neurons.

In conclusion, maintenance on a ketogenic diet does not inhibit voluntary activity on a running wheel, such that there are similar levels of activity in rats maintained on a ketogenic diet or chow. Whereas maintenance on a ketogenic diet leads to significantly increased adiposity in the rat, it is prevented with levels of exercise similar to those of chow-fed rats. The present results also indicate that voluntary exercise results in a significant increase in POMC expression in rats consuming a ketogenic diet, and highlight diet-related effects on the central melanocortin system that warrant attention for future study.

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Figure 1. Caloric intake in sedentary and exercising rats
Rats were maintained on CH or KD. Half of each group was allowed access to a running wheel (CH-RUN and KD-RUN), whereas the other half remained sedentary (CH-SED and KD-SED). Average caloric intake (kcal/day) during each week of the experiment is given for sedentary (SED) and exercising (RUN) rats that were maintained on CH or KD. At baseline, all rats were sedentary. Weeks 1–4 represent mean caloric intake during each of the 4 weeks the exercising rats had access to running wheel. Data are means ± SEM (kcal/day). *P < 0.05 when sedentary rats were compared to exercising rats.
Figure 2. Effects of access to a running wheel on body weight in chow- or ketogenic diet-fed rats
Body weights were also recorded daily for the duration of the experiment. After 2 weeks with access to a running wheel, rats on CH and rats on KD gained significantly less body weight than did sedentary controls. This difference persisted for the duration of the experiment. Data are means ± SEM body weight change (g) from the time access to the running wheel was allowed in the RUN groups. *$P < 0.05$ when RUN rats were compared to SED rats on the same diet.
Figure 3. Cumulative running wheel activity

Rats on CH or KD were allowed continuous access to running wheels attached to their home cages. (A) Total distance (m) over time and (B) rate of running (m/min) are given as mean ± SEM for CH or KD. There were no statistically significant differences between dietary groups for either measure.
Figure 4. Plasma analyses of sedentary and exercising rats

After 1 month during which rats on CH or KD were sedentary or allowed access to running wheels, plasma was analyzed by RIA for levels of (A) insulin and (B) leptin. Data are given as mean ± SEM and bars with different symbols are significantly different from one another (P < 0.05).
Figure 5. Hypothalamic neuropeptide mRNA expression in exercising and sedentary rats maintained on chow or a ketogenic diet

Neuropeptide mRNA expression levels were assessed by in situ hybridization for arcuate nucleus (A) NPY and (B) POMC. Data are means ± SEM and *P < 0.05 as compared to CH-SED.
### Table 1

Components of the ketogenic diet (Research Diets, D06040601).

| Ingredient                  | Gram | Kcal |
|-----------------------------|------|------|
| Protein                     | 23   | 15   |
| Carbohydrate                | 8    | 5    |
| Fat                         | 55   | 80   |
| Total                       |      | 100  |
| Kcal/g                      | 6.1  |      |

| Ingredient                  | gram | kcal |
|-----------------------------|------|------|
| Casein, 80 Mesh             | 151  | 604  |
| L-Cysteine                  | 1.5  | 6    |
| Corn Starch                 | 40   | 160  |
| Cellulose                   | 50   | 0    |
| Soybean Oil                 | 25   | 225  |
| Lard                        | 336  | 3024 |
| Mineral Mix, S10026         | 10   | 0    |
| Dicalcium Phosphate         | 13   | 0    |
| Calcium Carbonate           | 5.5  | 0    |
| Potassium Citrate, 1 H2O    | 16.5 | 0    |
| Vitamin Mix, V10001         | 10   | 40   |
| Choline Bitartrate          | 2    | 0    |
| FD&C Yellow Dye #5          | 0.025| 0    |
| FD&C Blue Dye #1            | 0.025| 0    |
| **TOTAL**                   | **660.55** | **4059** |

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Table 2

Effects of exercise and diet-type on cumulative caloric intake, body weight gain and adiposity in rats maintained on chow or a ketogenic diet. Values with differing symbols are significantly different from one another ($P < 0.05$).

|                          | CH-SED            | CH-RUN            | KD-SED            | KD-RUN            |
|--------------------------|-------------------|-------------------|-------------------|-------------------|
| Cumulative intake during | 3069.1 ± 95.0     | 2940.6 ± 82.0     | 2876.4 ± 100.7    | 2967.1 ± 36.3     |
| experiment (pre- and post-wheel kcals) |                   |                   |                   |                   |
| Cumulative intake with wheel access (kcals) | 2496.8 ± 84.4     | 2401.1 ± 64.3     | 2308.5 ± 86.5     | 2432.9 ± 27.3     |
| Terminal body weight (g) | 455.03 ± 16.01    | 412.78 ± 10.98*   | 442.73 ± 32.6     | 392.48 ± 15.43*   |
| Body weight gain from start of experiment (g) | 86.6 ± 4.00       | 67.68 ± 6.70*     | 88.27 ± 12.09     | 51.77 ± 8.17*     |
| Total fat pad weight (g) | 9.51 ± 1.11       | 6.93 ±0.96        | 16.66 ± 2.86*     | 8.82 ± 0.51       |
| Total fat pad (% body weight) | 2.18 ± 0.26       | 1.66 ± 0.18       | 3.78 ± 0.46*      | 2.26 ± 0.14       |
| Epidydimal fat (% body weight) | 0.76 ± 0.09       | 0.59 ± 0.07       | 1.23 ± 0.16*      | 0.75 ± 0.06       |
| Retroperitoneal fat (% body weight) | 0.73 ± 0.09       | 0.57 ± 0.09       | 1.31 ± 0.17*      | 0.84 ± 0.19       |
| Subcutaneous fat (% body weight) | 0.70 ± 0.08       | 0.49 ± 0.04*      | 1.24 ± 0.16#      | 0.67 ± 0.05       |