Resistance Exercise Increases the Capacity of Heme Biosynthesis More Than Aerobic Exercise in Rats

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Summary To evaluate the effects of voluntary resistance exercise and aerobic exercise on iron status in rats fed an iron-deficient diet (5 mg Fe/kg, ID) or rats fed a control diet (35 mg Fe/kg, CN), we trained female Wistar rats for 3 weeks to climb a wire-mesh tower (Ø 20 cm × 200 cm, CLIMB) and to swim in a plastic pool (Ø 50 cm × 50 cm, SWIM). These animals were compared with sedentary (SED) rats. After the experimental period, the blood hemoglobin concentration, hematocrit, plasma iron, and transferrin saturation were significantly lower (p<0.05) in the ID rats than in the CN rats; as was the content in the liver, spleen, heart, kidney, skeletal muscles, and carcass (p<0.05). The activity of δ-aminolevulinic acid dehydratase, the marker enzyme for heme biosynthesis, in bone marrow was significantly higher (p<0.05) in the CLIMB group than in the SWIM group. These results suggest that resistance exercise increases heme biosynthesis more than aerobic exercise but that neither exercise improved severe iron deficiencies.

Key Words: resistance exercise, swimming exercise, iron deficiency, δ-aminolevulinic acid dehydratase, heme biosynthesis

Iron deficiency continues to be a significant nutritional problem around in the world [1, 2]. It has deleterious effects on work performance, immune function, sympathetical and endocrinal metabolism, and thermoregulatory performance [3–6]. Although the hematologic and functional consequences of iron deficiency have been classically ascribed to a dietary or pathologic origin [2], previous studies suggest that chronic exercise might detrimentally alter body iron physiology.

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Decreased hematocrit, hemoglobin and serum iron, and increased erythrocyte fragility, may occur in aerobic exercising individuals [7-10]. Thus far, animal studies [11, 12] that examined iron deficiency and aerobic exercise interactions demonstrated that exercise lessens the impact of moderate iron deficiency on essential body iron components, such as hemoglobin. Hisaoka and Shibuya [13] demonstrated that hemoglobin, hematocrit, and blood red cell volume were significantly higher in female swimming rats than in control rats. Tobin and Beard [14] reported that running failed to alter hemoglobin, hematocrit and red blood cell mass in iron-deficient and control rats.

On the other hand, studies on the effects of resistance exercise on body iron status are limited in number. We previously demonstrated that mild resistance exercise improved non-anemic iron deficiency without iron supplementation in young women [15]. However, the mechanism of the effects of resistance exercise on iron metabolism is not clear.

Recently, we designed a new voluntary resistance training model, in which rats climbed a vertical tower [16]. The purpose of the present study was to determine whether resistance exercise induces an improvement of essential body iron components in iron-deficient rats and, if so, to determine how the increase in δ-aminolevulinic acid dehydratase (ALAD) activity in bone marrow, a marker enzyme of mitochondrial heme biosynthesis, compares with that found with aerobic exercise.

**MATERIALS AND METHODS**

All procedures involving animals were approved by the Experimental Animal Care Committee of Kagawa University.

**Animals and experimental design.** Twenty-four female Wistar rats (3 weeks old) were obtained from Japan SLC, Inc. (Shizuoka, Japan). The rats were fed CE-2, commercial rodent diet (CLEA Japan, Tokyo), and given water *ad libitum* through 4 weeks of age. All animals were individually housed in an animal room at 24±1°C, with lights on from 8 a.m. to 8 p.m. Half of the animals were assigned to the AIN-76 diet [17, 18], with less than 5 mg Fe/kg (iron deficiency, ID), and the other half were assigned to an identical diet with 35 mg Fe/kg (control, CN). The ID and CN rats were each randomly divided into 3 subgroups, sedentary (SED), swimming exercise (SWIM), and climbing exercise (CLIMB) groups. Each group of rats (n=4/group) was given free access to the ID or CN diet, and water for 21 days. After the 21-day experimental period, the rats were fasted overnight and killed by decapitation at 10 a.m. under light ether anesthesia. Liver, spleen, heart, kidney, and skeletal muscles were quickly removed and stored at −40°C until analysis could be performed. Carcass samples were obtained by removing the head, digestive tracts, lungs, testes, and abdominal adipose tissues, and stored at −20°C until analyzed.

**Exercise training.** The voluntary resistance training model, CLIMB [16],
was a modification of that described by Yarasheki et al. [19] and Duncan et al. [20]. Rats of the CLIMB group were housed in metal cages containing wire-mesh towers (φ 20 cm × 200 cm) with water bottles set on the top of the tower [16]. On the other hand, the rats of the SWIM group were trained from 8 a.m. to 9 a.m. every day in a plastic pool (φ 50-50 cm) with water maintained at 33–35°C. The swimming exercise was performed in the manner described previously [21, 22]. All rats were rested for 24 h in their individual cages before sacrifice.

**Blood and plasma analysis.** Blood hemoglobin concentrations were determined colorimetrically by using a hemoglobin B-Test kit purchased from Wako Pure Chemical Industries (Osaka, Japan). The hematocrit was measured by centrifugation of blood collected into heparinized microcapillary tubes. Plasma iron concentration and total iron binding capacity (TIBC) were determined by the method prescribed by the International Nutritional Anemia Consultative Group [23]. Transferrin saturation was calculated by plasma iron concentration and TIBC [21].

**Tissue and carcass iron.** Carcass samples were prepared by the method of Mickelsen and Anderson [24]. Iron contents in the liver, spleen, heart, kidney, soleus muscle, white and red gastrocnemius muscles, and carcass were measured after acid hydrolysis by using an atomic absorption spectrophotometer (Model Z-5000, Hitachi, Tokyo, Japan).

**Enzyme assay.** ALAD activity in the bone marrow was measured by the method of Sassa [25], with the modification that Ehrlich's reagent was made as reported by Tomokuni [26]. Briefly, about 30 mg of bone marrow was obtained from the femur and homogenized in a micro homogenizer (ULTRA-TURRAX, IKA Co., Staufen, Germany) at 0-4°C containing 270 μl of 0.1 M potassium phosphate buffer (pH 6.4). Part of the homogenate (100 μl) was transferred to a 1.5-ml tube, and 250 μl of 0.1 M potassium phosphate buffer containing 10 mM δ-aminolevulinic acid was added. After incubation for 30 min at 37°C, the reaction was terminated by the addition of 250 μl of 10% trichloroacetic acid. The precipitate was centrifuged at 15,000 × g for 10 min. A 0.5-ml volume of the supernatant was removed into another tube, and 0.5 ml of modified Ehrlich's reagent was added. After 40-60 min, the color solution absorbance was determined at 553 nm vs. the blank containing 0.5 ml of 0.1 M potassium phosphate buffer and 0.5 ml of modified Ehrlich's reagent. The ALAD activity was calculated as follow: absorbance × 2 × 120 × 2/6.2 (μmol/h/g tissue) (2, conversion factor from porphobilinogen to δ-aminolevulinic acid ; 120, dilution factor ; 2, incubation time ; 6.2, extinction coefficient in ml/μmol·cm). Protein content in the reaction mixture was determined by the method reported by Lowry et al. [27].

The activities of citrate synthase in the heart and plantaris muscle were determined by the method described by Srere [28] to establish whether the training protocol had any effect on aerobic metabolism. One unit of the citrate synthase catalyzed the formation of 1 μmol free CoA/min [28].

**Statistics.** Data were expressed as means ± SE. All data were analyzed by a
factorial analysis of variance (ANOVA) and Fisher’s PLSD tests. Differences were considered statistically significant at $p<0.05$.

RESULTS

Body weight, food intake, and food efficiency

Body weight changes over 21 days and body weight gain are shown in Fig. 1. Exercise training lowered body weight gain in both ID and CN rats. In ID rats, the body weight gain was significantly lower ($p<0.05$) in the CLIMB group than in the SED group (Fig. 1). Food intake was approximately the same in all experimental groups (284±12, 271±8, 296±4, 270±8, 273±10, and 280±2 g/21 days for CN-SED, CN-SWIM, CN-CLIMB, ID-SED, ID-SWIM, and ID-CLIMB groups, respectively). Food efficiency was significantly lower ($p<0.05$) in the SWIM and CLIMB groups than in the SED group for both ID and CN rats (186±5, 178±1, 166±3, 188±7, 168±6, and 153±3 mg body weight/g for CN-SED, CN-SWIM, CN-CLIMB, ID-SED, ID-SWIM, and ID-CLIMB groups, respectively).

Hemoglobin, hematocrit, plasma iron, and transferrin saturation

The results of hemoglobin, hematocrit, plasma iron, and transferrin saturation measurements are shown in Table 1. All components were significantly lower ($p<0.05$) in the ID rats than in the CN rats for SED, SWIM, and CLIMB groups. However, neither exercise affected the iron status of the ID or CN rats.

![Fig. 1. Body weight changes over 21 days and weight gains of sedentary and exercised rats. Values are means and SE (weight gain graph only) for 4 rats. *Statistically significant difference from the sedentary group ($p<0.05$, ANOVA and Fisher’s PLSD tests). ANOVA indicated significant ($p<0.05$) main effects of exercise.](image)
Tissue and carcass iron contents

The iron content in the liver, spleen, heart, kidney, soleus muscle, red and white gastrocnemius muscle, and carcass was significantly lower ($p < 0.05$) in the ID rats than in the CN rats (Table 2). However, neither exercise affected the iron status of the ID or CN rats except for the soleus iron content in the ID-CLIMB group (Table 2). Liver iron was more strongly affected by the ID diet (67%
Enzyme activity

The climbing exercise increased but swimming exercise did not increase, bone marrow ALAD activity in both ID and CN groups (Fig. 2). The bone marrow ALAD activity was significantly higher ($p<0.05$) in the CLIMB group than in the SED and SWIM groups (Fig. 2). Citrate synthase activities in both plantaris muscle and heart were increased by exercise training (Table 3). Climbing exercise decreases for both SWIM and CLIMB groups) than the iron in other tissues.

**Table 3. Effects of exercise and iron deficiency on citrate synthase activity in plantaris muscle and heart of rats**

<table>
<thead>
<tr>
<th>Group</th>
<th>$n$</th>
<th>Plantaris (U/g tissue)</th>
<th>Heart (U/g tissue)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CN-SED</td>
<td>4</td>
<td>2.9 ± 0.5</td>
<td>3.21 ± 0.4</td>
</tr>
<tr>
<td>CN-SWIM</td>
<td>4</td>
<td>2.8 ± 1.0</td>
<td>10.5 ± 1.1</td>
</tr>
<tr>
<td>CN-CLIMB</td>
<td>4</td>
<td>6.7 ± 0.9 $^{#}$</td>
<td>9.4 ± 1.3</td>
</tr>
<tr>
<td>ID-SED</td>
<td>4</td>
<td>2.4 ± 0.9</td>
<td>5.5 ± 1.6</td>
</tr>
<tr>
<td>ID-SWIM</td>
<td>4</td>
<td>3.4 ± 0.9</td>
<td>7.5 ± 0.4</td>
</tr>
<tr>
<td>ID-CLIMB</td>
<td>4</td>
<td>4.9 ± 0.8 $^{#}$</td>
<td>10.7 ± 1.4</td>
</tr>
</tbody>
</table>

$^{a}$Values are means ± SE for 4 rats. Statistically significant difference from the sedentary. $^{b}$Statistically significant difference from the swimming exercise group ($p<0.05$, ANOVA and Fisher's PLSD tests). ANOVA indicated significant ($p<0.05$) main effects of exercise for all variables. $^{c}$CN, control; ID, iron deficiency; SED, sedentary; SWIM, swimming exercise; CLIMB, climbing exercise.
by rats resulted in a significant increase \( (p<0.05) \) in citrate synthase activity in both the plantaris muscle (104 and 131% for ID and CN groups, respectively) and heart (95 and 193% for ID and CN groups, respectively). Swimming exercise significantly \( (p<0.05) \) increased the activity of heart citrate synthase in the CN rats, but did not activate plantaris citrate synthase (Table 3).

DISCUSSION

This study demonstrated that resistance exercise (CLIMB) increased the capacity of heme biosynthesis in the bone marrow more than did aerobic exercise (SWIM). These results suggest that the capacity for heme biosynthesis may be dependent on resistant loads to the animal body. Considerable evidence indicates that ALAD, the enzyme in the porphyrin pathway, is a marker enzyme for heme biosynthesis \[29\]. Thus, increases in the activity of this enzyme in the CLIMB rat bone marrow have been linked to accelerated heme production. Hemoglobin is dependent on heme synthesis in the bone marrow since iron-containing porphyrin constitutes the ring structure to which the hemoglobin is conjugated with apoprotein. Therefore, we considered it likely that the heme pathway would be accelerated when exercise-induced increases in the hemoglobin concentration could be expected. However, our results showed no exercise-induced effects on the capacity for heme biosynthesis as reflected in hemoglobin production as well as on other blood components. Tobin and Beard \[14\] suggested that training in iron-deficient animals was characterized by a higher percentage of \(^{59}\)Fe associated with red cells than that for iron-deficient sedentary rats within 1 h after intravenous injection of \(^{59}\)Fe. But they failed to show any improvement in the blood hemoglobin level in iron-deficient trained rats. Franzone et al. \[30\] determined that the blood red cell mass and hemoglobin were the initial sites of plasma iron clearance. A reduction in plasma iron circulation caused by severe iron deficiency may void exercise-induced alterations in the capacity for heme biosynthesis.

In this study, the iron content in several tissues was significantly lower in the ID rats, and neither exercise affected the iron status of the ID or CN rats except for the soleus iron content in ID-CLIMB group. Liver iron was more severely reduced by the ID diet than the iron in other tissues. These findings are consistent with the fact that the liver is the main control site for plasma iron circulation \[31\]. Borel et al. \[32\] reported that as the dietary iron intake was increased above 11 mg Fe/kg diet, the liver iron concentration steadily increased, and the hemoglobin concentration was maintained at normal levels. Siimes et al. \[33\] determined that a dietary iron intake of less than 25 mg Fe/kg diet resulted in hemoglobin concentrations of 12 mg/100 ml or below. From these points of view, the dietary iron level in the ID diet (less than 5 mg Fe/kg diet) that we used in this study might have been too low to examine the effects of exercise on body iron status.

Previous studies \[34, 35\] showed that iron deficiency did not suppress animal growth during a 12-week experimental period. We also failed to observe a signifi-
cant depression in body weight attributable to iron deficiency. However, both exercises depressed body weight gain in both dietary treatment groups, with the iron-deficient CLIMB rats showing the greatest attenuation of growth (12.8%). On the other hand, food intake was at the same level in all experimental groups. These data demonstrate that fat mass is reduced in ID-CLIMB rats compared to ID-SED rats, since muscle mass and carcass protein content did not differ among the 6 groups (data not shown). Tobin and Beard [14] reported that both iron deficiency and running significantly increased resting oxygen consumption, as an index of resting energy expenditure, 48 h postexercise in rats. Consequently, fat mass was significantly lower in iron-deficient exercised rats than in iron-deficient sedentary rats. Our present study supports the results of Tobin and Beard.

Citrate synthase activities in the plantaris muscle and heart were increased by exercise training. CLIMB training by rats resulted in a significant increase in citrate synthase activity in both the plantaris muscle and heart. These results are in agreement with our previous findings [16]. On the other hand, swimming significantly increased the activity of heart citrate synthase in the CN rats, but did not activate plantaris citrate synthase. Voluntary swimming exercise may increase the aerobic capacity of the heart prior to that of the skeletal muscles.

In conclusion, our study suggests that resistance exercise increases the capacity of heme biosynthesis in the bone marrow as compared with aerobic exercise. However, resistance exercise did not improve severe iron deficiencies. Further study using rats at other iron-deficiency levels will be required to clarify the details concerning the effects of resistance exercise on iron-deficiency anemia.

REFERENCES
