Blood Glucose Regulation during Fasting in Rats under Food Restriction since Birth

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ABSTRACT

The effect of severe food restriction since birth on regulation of fasting glycemia in male Wistar rats was investigated. The control group (CG) had free supply of chow, while the restriction group (RG) received 50% of the amount ingested by the CG. The experiments were done in adult (60 days) overnight fasted rats in which glycemia, liver free glucose levels and hepatic glycogen concentration were measured. In part of the experiments in situ liver perfusion was done. The results showed that livers from the RG had higher glycogenolysis rates but lower gluconeogenesis rates from L-alanine (10 mM). Since RG showed maintained glycemia during fasting, it could be concluded that livers from RG produced glucose preferentially from glycogenolysis in detriment of gluconeogenesis. These findings demonstrated that in spite of severe caloric restriction, the metabolic adaptations of the liver did exist to assure the maintenance of blood glucose for brain supply during fasting.

Key words: food restriction, rats, liver, gluconeogenesis, glycogenolysis

INTRODUCTION

Undernutrition in children is characterized by lower physical growth and mental development (De Onis and Blossner, 2003; FAO, 2003; WHO, 2005). However, in rats, undernutrition caused by food restriction since birth has marked effects on body weight development, accompanied by reduction in the muscle mass and fat stores (Mazeti and Furlan, 2008; Vismara and Furlan, 2007). Several studies have demonstrated changes in the liver metabolism in adult rats submitted to food restriction (Batista et al., 2001; Bazotte et al., 1989, 2000). These alterations include changes in liver gluconeogenesis, nitrogen removal, and activation of the transaminases (Dhahbi et al., 1999; Hagopian et al., 2003; Spindler, 2001). However, information about metabolic changes in rats submitted to food restriction since birth, particularly about the blood regulation of glycemia during fasting, are scarce. Since L-alanine (L-Ala) is the most important liver glucose precursor in conditions of hypoproteic diet or prolonged fasting (Brosnan et al., 2001), we investigated if severe food restriction since birth could influence the liver metabolism of L-Ala. Moreover, the liver

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glycogenolysis and several other metabolic parameters were evaluated.

MATERIALS AND METHODS

Animals
Pregnant Wistar rats were housed in individual cages, where they gave birth about a week after their arrival. The newborn litters were reorganized so that each mother had either six (control group, CG) or 12 pups (restriction group, RG). Only male animals were used in this investigation. The dams had free access to water and food (Nuvilab CR1®; Nuvital, Curitiba, Brazil) during gestation and lactation. The pups remained with their mothers until the age of 21 days (weaning), when they were put in groups of four or five in plastic boxes with free access to water. The CG rats also had free supply of chow, while those in the RG group were subjected to a 50% food restriction (Mazeti and Furlan, 2008; Vismara and Furlan, 2007). No dietary supplementation was given to the RG. All the experiments were done after overnight fasting when the animals were 45-60 days of age. The rats were kept under constant light:dark cycles (12 h light:12 h dark) and temperature (22 ± 2°C). All the procedures were approved by the Committee of Ethical Conduct in the Use of Animals in Experimentation (CEAE) of the State University of Maringá.

Collection of blood and tissues
The rats from CG and RG groups were weighed and the blood to measure glycemia was collected from the caudal vein. After that the rats were killed by decapitation and the liver and the retroperitoneal and periepididymal fat pads (left and right) were removed and weighted. The blood collected in heparinized tubes was centrifuged (3,000 rpm for 5 min) and the plasma was separated to determine glycemia (Bergmeyer and Bernt, 1974).

Hepatic free glucose and glycogen content
Livers from CG and RG rats were quickly removed, immersed in liquid nitrogen and weighted. About 1.5 g of the organ was grinded while kept frozen by the addition of liquid nitrogen. Perchloric acid (0.6 N) was added and the resulting mass was homogenized and centrifuged (10 min at 6,000 rpm). Aliquots of 100 µL of the supernatant were used to determine the levels of free glucose (Bergmeyer and Bernt, 1974). To another aliquot of supernatant (100 µL), amylglucosidase (50 µL) was added along with potassium bicarbonate (50 µL) and sodium acetate (960 µL). The solution was incubated at 40°C in water bath under agitation for two hours, and the enzymatic reaction was interrupted by the addition of perchloric acid (0.6 N, 500 µL). Finally, after another centrifugation (10 min at 6,000 rpm), the aliquots of supernatant (100 µL) were used to determine total glucose concentration, i.e., free glucose plus glycogen-derived glucose (Bergmeyer and Bernt, 1974; Oliveira et al., 2007).

In situ liver perfusion
The rats were anesthetized with an intraperitoneal injection of sodium thiopental (40 mg/kg bw) and submitted to laparotomy. The livers were perfused in situ using Krebs Henseleit bicarbonate buffer (pH 7.4), saturated with O2/CO2 (95/5%). The perfusion fluid was pumped through a temperature-controlled (37°C) membrane oxygenator prior to entering the liver via the portal vein. Samples of the effluent perfusion fluid were collected from the inferior cava vein each 5 min and the concentrations of glucose, pyruvate (Czok and Lamprecht, 1974), L-lactate (Gutmann and Wahlefeld, 1974) and urea (Gutmann and Bergmeyer, 1974) were measured. As shown in Figure 1, after a pre-infusion period (15 min without L-Ala), L-Ala (10 mM) was dissolved in the perfusion fluid and infused during 20 min, followed by a post-infusion period (10 min without L-Ala). The glucose production before the infusion of L-Ala represented the glucose from glycogenolysis and the difference in the production of glucose during and before the infusion of L-ala represented
the rate of gluconeogenesis. During the infusion of L-al, along with glucose, the production of pyruvate, L-lactate and urea was also measured. The differences in the production of glucose, pyruvate, L-lactate and urea during and before the infusion of L-Ala allowed to calculate the area under the curves (AUC), expressed as µmol/g liver (Table 3).

![Graph](image)

**Figure 1** - Demonstrative experiment of a perfusion protocol showing glucose production before (pre-infusion period), during and after (post-infusion period) the infusion of L-alanine 10 mM (L-Ala).

### Statistical analysis

The data were statistically treated using non-paired Student’s t test. P values < 0.05 were considered significant. The statistical analyses were carried out with Prism v4. The data are shown as means (M) ± standard deviations (SD).

### RESULTS AND DISCUSSION

Rats from the RG group showed lower (P < 0.05) body weight from lactation until adult age (Table 1). Therefore, when food restriction since birth was given, the body weight gain was lower, as previously described (Aubert et al., 1980; Duff and Snell, 1982; Houdijk et al., 2003). This could be explained by the limited food supply which was not enough to meet the caloric requirement of the RG group (Emery, 2005; Mazeti and Furlan, 2008; Vismara and Furlan, 2007). The RG showed lower (P < 0.05) relative weights of retroperitoneal and periepididymal fat pads (Table 2). This has been observed when nutrients are available in amounts lower than those needed to secure a normal body growth (Barzilai et al., 1999; Dulloo and Girardier, 1993; Porter et al., 2004; Valle et al., 2005; Waterland and Garza, 1999).

<table>
<thead>
<tr>
<th>Age</th>
<th>CG (n=5-8)</th>
<th>RG (n=5-8)</th>
</tr>
</thead>
<tbody>
<tr>
<td>5 days</td>
<td>10.25±1.17</td>
<td>9.67±0.82</td>
</tr>
<tr>
<td>20 days</td>
<td>36.67±3.78</td>
<td>27.50±2.03*</td>
</tr>
<tr>
<td>60 days</td>
<td>245.21±4.31</td>
<td>131.58±11.05*</td>
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Table 1 - Body weight (g) of rats with free access to food (CG) and rats submitted to food restriction (RG) since birth. *P < 0.01 relative to the CG of the same age; non-paired Student’s t test. n = number of rats.

<table>
<thead>
<tr>
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<th>CG (n=5-8)</th>
<th>RG (n=5-8)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Liver (g/100 g body weight)</td>
<td>3.00±0.22</td>
<td>3.93±0.46*</td>
</tr>
<tr>
<td>Retroperitoneal fat pads (g/100 g body weight)</td>
<td>1.14±0.22</td>
<td>0.24±0.11*</td>
</tr>
<tr>
<td>Periepididymal fat pads (g/100 g body weight)</td>
<td>0.99±0.18</td>
<td>0.51±0.11*</td>
</tr>
<tr>
<td>Liver caloric density (kJ/g dry weight)</td>
<td>23.54±0.11</td>
<td>21.16±0.51*</td>
</tr>
</tbody>
</table>

Table 2 - Relative weight of the liver, retroperitoneal and periepididymal fat pads and caloric density of the liver in the control group (CG) and restriction group (RG). *P < 0.05 relative to the CG; non-paired Student’s t test. n = number of animals.
In contrast, the liver weight was higher (P < 0.01) in the RG, indicating that the liver had priority when the nutritional supply was not enough to furnish the metabolic energy for all tissues and organs (Porter et al., 2004). Calorimetry is a method that provides the amount of total calories from protein, carbohydrate or fat of a dehydrated tissue. The present results showed lower (P < 0.05) caloric density in livers from RG. Since lipids have higher caloric content than protein and carbohydrates, the decreased caloric density in the RG could be the result of a reduction of the lipid content of the liver. Thus, the reduced liver caloric content and the reduced fat pad weight pointed to a general decrease in the total content of body fat.

The total hepatic content of glucose, i.e., free glucose plus glycogen-derived glucose, did not differ (294.1±5.45 and 297.6±4.95 µmol/g liver in the CG and RG groups, respectively), but the content of free glucose was higher (P < 0.01) in the RG than the CG (5.75±1.86 and 3.07±1.62 µmol/g liver, respectively). In agreement, the basal hepatic glucose production, i.e., before L-alanine infusion, was higher (P < 0.05) in the livers from RG.

Considering that the glucose production before L-Ala infusion resulted from glycogen (Batista et al., 2001), it could be inferred that the livers from RG showed a higher rate of glycogenolysis despite the fact that the hepatic glycogen content was not significantly different between the groups.

In contrast, lower (P < 0.05) liver production of glucose and L-lactate during the infusion of L-Ala was observed. However, the production of urea and pyruvate were similar (Table 3).

Table 3 - Glucose production before (pre-infusion period, 10 min) and during (20 min) the infusion of L-Ala (10 mM) in livers from control (CG) and restriction groups (RG). During the infusion of L-alanine, the liver production of glucose, pyruvate, L-lactate and urea were measured. *P < 0.05 relative to the CG; non-paired Student’s t test. n = number of animals.

<table>
<thead>
<tr>
<th>Liver production (µmol/g liver)</th>
<th>CG (n=5-8)</th>
<th>RG (n=5-8)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Before L-alanine infusion (Basal perfusion)</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glucose</td>
<td>0.63±0.17</td>
<td>3.34±1.20*</td>
</tr>
<tr>
<td><strong>During L-alanine infusion</strong></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Glucose</td>
<td>0.83±0.17</td>
<td>0.45±0.17*</td>
</tr>
<tr>
<td>Pyruvate</td>
<td>1.74±0.04</td>
<td>2.12±0.57</td>
</tr>
<tr>
<td>L-Lactate</td>
<td>1.34±0.42</td>
<td>2.10±0.63*</td>
</tr>
<tr>
<td>Urea</td>
<td>2.37±0.42</td>
<td>2.40±0.55</td>
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</table>

Considering that the increased glucose production during L-Ala infusion resulted from gluconeogenesis (Galende et al., 2009), it could be inferred that the livers from RG showed lower rate of gluconeogenesis from L-Ala. In agreement with this result, Liu et al. (1996) showed reduced liver uptake and lower gluconeogenesis from L-Ala in the rats under food restriction. Hagopian et al. (2003) demonstrated that the carbon skeleton of amino acids could be shifted to ketogenesis during food restriction, and in long-term food-restricted rats, ketogenesis is highly favored (Emmanuel and Kennely, 1984; Field, 1989; Westman et al., 2007). Therefore, it was possible that the nitrogen balance in the RG could be funneling L-Ala to other compounds by transamination as a consequence of the reduced protein supply in the diet. In agreement, increased hepatic transaminases and turnover of extra-hepatic proteins during food restriction were reported (Hagopian et al., 2003; Spindler, 2001).

Finally, the main result of this study was the demonstration that food restricted rats since birth showed higher glycogenolysis and lower gluconeogenesis. Thus, considering that gluconeogenesis is a metabolic pathway with high energetic cost, it could be suggested that the preference for glycogenolysis could be an adaptation for glycemia maintenance during fasting to overcome the lower food availability (Batista et al., 2001; Bazotte et al., 2000). In conformity with this suggestion, fasting glycemia did not differ (P > 0.05) between CG (97.3±23.3 mg/dL) and RG groups (103.3±25.9 mg/dL).
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Received: April 20, 2010; Revised: July 29, 2010; Accepted: November 17, 2010.