Erythrocyte Glycolysis and Its Marked Alteration by Muscular Exercise in Type VII Glycogenosis

By Takao Shimizu, Norio Kono, Hiroaki Kiyokawa, Yuya Yamada, Naoko Hara, Ikuo Mineo, Masanori Kawachi, Hiromu Nakajima, Yan Lin Wang, and Seiichiro Tarui

Levels of erythrocyte glycolytic intermediates after the phosphofructokinase (PFK) step, including 2,3-bisphosphoglycerate (2,3-DPG), were decreased at rest in patients from separate families with type VII glycogenosis. The concentration of 2,3-DPG was about half of the normal control value during a period of unrestricted daily activity but was further decreased to one third of normal after a one-day bed rest. Mild ergometric exercise rapidly increased the levels of fructose-1,6-bisphosphate, dihydroxyacetone phosphate plus glyceraldehyde-3-phosphate, and 2,3-DPG in patients' circulating erythrocytes but did not in those of normal subjects. This indicated that a crossover point at the PFK step in glycolysis disappeared after physical exercise and, consequently, the 2,3-DPG concentration, which had decreased because of blockage of the PFK step, was restored considerably. This apparently exercise-related alteration in intermediary metabolism at the beginning of glycolysis was reproduced in vitro by incubating normal erythrocytes in the presence of inosine or ammonia, both of which have increased levels in circulating blood during and after exercise in this disorder. We conclude that physical activity in addition to a genetic deficiency in erythrocyte PFK affects glycolysis in erythrocytes in type VII glycogenosis and that myogenic factors released from exercising muscles may be responsible for this change.

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In type VII glycogenosis (muscle and erythrocyte phosphofructokinase [PFK] deficiency), the activity of erythrocyte PFK is reduced to about half of the normal value. Erythrocyte 2,3-bisphosphoglycerate (2,3-DPG) is decreased because of blockage of erythrocyte glycolysis at the PFK step. Recently we noticed that the levels of erythrocyte 2,3-DPG fluctuated and that this alteration seemed to be related to the patients' physical activity. Moreover, we have reported that large quantities of ammonia, purine nucleosides, and oxypurine are released into the blood from exercising muscles in type VII glycogenosis. Ammonia is known to be a potent activator of PFK, and inosine is readily metabolized in erythrocytes through the hexose monophosphate shunt. Therefore, the intermediary metabolism of glycolysis in circulating erythrocytes might be affected by the increase in these compounds in blood during and after exercise in patients with type VII glycogenosis. In this study, we have examined the effect of muscular exercise and bed rest on the glycolysis of circulating erythrocytes in type VII glycogenosis.

MATERIALS AND METHODS

Patients. Erythrocytes from four patients with type VII glycogenosis were the subjects of this study. Patient 1 (a 46-year-old male) and patient 2 (a 43-year-old male) belong to the same family reported in 1965 as the probands of this disease. Patient 3 (a 19-year-old male) and patient 4 (a 16-year-old female) belong to another family in Japan. The kinetics of residual erythrocyte and liver PFK in all patients were similar. All the blood samples from the patients were drawn directly into a syringe containing 6% perchloric acid. Patient 2 exercised only intermittently on a bicycle ergometer because of exercise intolerance. The subject completed two 35-W work loads in five minutes with a ten-minute rest and then a 40-W load in eight minutes. Erythrocytes drawn from patient 2 after the bicycle ergometer exercise were also analyzed. Procedures and consent forms were approved by the Osaka University Hospital Committees on the Protection of Human Subjects in Research.

Biochemical studies. Glycolytic intermediates were assayed as described previously. The plasma ammonia concentration was measured enzymatically. Inosine was measured by a high-performance liquid chromatography system (Gilson) with a microsorb C18 reversed-phase column (Rainin Instrument Co, Woburn, MA). The adenylic acid energy charge was calculated by the formula given by Atkinson and Walton.

An erythrocyte incubation study was performed as described previously. Leukocytes and platelets were removed from heparinized blood by filtration through a cellulose column. Then the erythrocytes were washed with 0.9% NaCl three times and suspended in a solution containing 100 mmol/L NaCl, 10 mmol/L KCl, 50 mmol/L triethanolamine/HCl, 30 mmol/L glucose, and 1 mmol/L inorganic phosphate (pH 7.4). The hematocrit of the suspension was 44%. After a ten-minute preincubation at 37°C, varying concentrations of ammonium chloride or inosine were added. The reaction was stopped by the addition of perchloric acid (final concentration, 3%) after 20 minutes.

All enzymes used in this study were purchased from Boehringer Mannheim and other reagents from Sigma Chemical Co (St Louis).

RESULTS

Erythrocyte metabolite concentrations at rest. The concentration of glycolytic intermediates and adenine nucleotides in erythrocytes from patients with type VII glycogenosis at rest are shown in Table 1. There were decreases in fructose-1,6-bisphosphate, dihydroxyacetone phosphate plus glyceraldehyde-3-phosphate, and 2,3-DPG in all three patients examined at rest during the period of unrestricted
Table 1. Erythrocyte Glycolytic Intermediates and Adenine Nucleotides in Type VII Glycogenosis

<table>
<thead>
<tr>
<th>Intermediate/ Nucleotide</th>
<th>Normal Value (Mean ± SD)</th>
<th>During Unrestricted Daily Activity</th>
<th>During Bed Rest</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Patient 1</td>
<td>Patient 2</td>
</tr>
<tr>
<td>G6P</td>
<td>35.6 ± 6.2</td>
<td>44.5</td>
<td>38.4</td>
</tr>
<tr>
<td>F6P</td>
<td>13.4 ± 3.1</td>
<td>13.9</td>
<td>12.2</td>
</tr>
<tr>
<td>FBP</td>
<td>1.73 ± 0.53</td>
<td>0.92</td>
<td>0.70</td>
</tr>
<tr>
<td>DHAP + GAP</td>
<td>15.6 ± 4.4</td>
<td>14.7</td>
<td>8.2</td>
</tr>
<tr>
<td>2,3-DPG</td>
<td>4,810 ± 550</td>
<td>2,950</td>
<td>2,430</td>
</tr>
<tr>
<td>3PGA</td>
<td>42.4 ± 4.6</td>
<td>50.7</td>
<td>22.7</td>
</tr>
<tr>
<td>2PGA</td>
<td>6.5 ± 1.9</td>
<td>5.8</td>
<td>7.1</td>
</tr>
<tr>
<td>PEP</td>
<td>13.8 ± 4.0</td>
<td>14.5</td>
<td>15.5</td>
</tr>
<tr>
<td>PYR</td>
<td>98.2 ± 28.0</td>
<td>56.1</td>
<td>61.6</td>
</tr>
<tr>
<td>LAC</td>
<td>911 ± 194</td>
<td>605</td>
<td>597</td>
</tr>
<tr>
<td>ATP</td>
<td>1,630 ± 132</td>
<td>1,040</td>
<td>1,170</td>
</tr>
<tr>
<td>ADP</td>
<td>203 ± 48</td>
<td>243</td>
<td>254</td>
</tr>
<tr>
<td>AMP</td>
<td>22.3 ± 8.6</td>
<td>36.8</td>
<td>31.7</td>
</tr>
<tr>
<td>Total adenylates</td>
<td>1,855 ± 172</td>
<td>1,320</td>
<td>1,456</td>
</tr>
<tr>
<td>Energy charge</td>
<td>0.935 ± 0.012</td>
<td>0.880</td>
<td>0.891</td>
</tr>
</tbody>
</table>

Values are expressed as μmol/L RBCs.
Abbreviations: G6P, glucose-6-phosphate; F6P, fructose-6-phosphate; FBP, fructose-1,6-bisphosphate; DHAP, dihydroxyacetone phosphate; GAP, glyceraldehyde-3-phosphate; 3PGA, 3-phosphoglycerate; 2PGA, 2-phosphoglycerate; PEP, phosphoenolpyruvate; PYR, pyruvate; LAC, lactate.
*Values in μmol/L blood.

Effect of exercise on erythrocyte metabolism. The effect of ergometric exercise on the glycolytic metabolism of erythrocytes in the systemic circulation was studied in a patient with type VII glycogenosis (patient 2). Immediately after exercise, concentrations of intermediates in the beginning of glycolysis were much increased, and those at the end became normal (Fig 3), whereas no significant change was found in normals (data not shown). The concentrations of

Fig 1. Crossover plot of the mean concentrations of glycolytic intermediates and adenine nucleotides in erythrocytes from patients with type VII glycogenosis. The values plotted were represented in terms of the ratios to control values. Data from Table 1 were used to construct the curve. *P < .05; **P < .01; and ***P < .001 vs controls.
fructose-1,6-bisphosphate and dihydroxyacetone phosphate plus glyceraldehyde-3-phosphate were about fourfold of control values. The AMP concentration became normal, and 2,3-DPG and ATP levels tended to increase during ergometric exercise. One hour after exercise, the concentrations of intermediates decreased but were still slightly higher than preexercise concentrations.

Although exercising was difficult for this patient, there was a significant increase in the 2,3-DPG concentration from 1,710 to 2,030 μmol/L RBC, but its concentration was not restored completely (Fig 2). To see the effect of prolonged but less intensive exercise on the 2,3-DPG concentration, 2,3-DPG was measured after the patient was discharged and had returned to work (the results are shown in Fig 2). Levels of 2,3-DPG increased from 2,010 to 2,420 μmol/L RBC, which is consistent with the finding that the 2,3-DPG concentration is decreased by prolonged rest.

**In vitro studies.** We previously showed that exercise on a bicycle ergometer markedly increased plasma inosine and hypoxanthine levels in this patient. The inosine concentration rose from 0.4 to 9.5 μmol/L and the ammonia concentration also rose from 60 to 475 μmol/L after exercise but was unchanged in normal subjects. To ascertain whether the exercise-related alteration of erythrocyte glycolysis in type VII glycogenosis is caused or mediated by the augmentation of inosine or ammonia, the effects of these metabolites on the intermediary metabolism of erythrocyte glycolysis were studied in vitro experiments, which are shown in Fig 4. The addition of inosine increased the levels of fructose-1,6-bisphosphate and dihydroxyacetone phosphate plus glyceraldehyde-3-phosphate. The increments of these intermediates were greater as the inosine concentration was increased up to 150 μmol/L. With the addition of 150 μmol/L inosine, 2,3-DPG levels increased from 4,280 to 4,730 μmol/L RBC in 20 minutes. The incremental amount of 2,3-DPG and lactate was more than the decremental amount of inosine, which may be in part due to the further activation of PKF by increased fructose-1,6-bisphosphate levels. The addition of ammonium chloride to give a final concentration of 1,000 μmol/L produced a forward crossover between fructose-6-phosphate and fructose-1,6-bisphosphate, thus indicating the increased flux at the PKF step.

**DISCUSSION**

Concentrations of erythrocyte glycolytic intermediates after the PKF step were decreased in two families with type VII glycogenosis (Table 1). This implies that the metabolic flux is decreased at the PKF step in vivo. The concentration of 2,3-DPG was about half that of normal during unrestricted daily activity and one third of that after bed rest. These changes in 2,3-DPG concentrations during different physical activities were observed in both families. Because erythrocyte PKFs of the patients from these families were

![Fig 2. Fluctuations of erythrocyte 2,3-DPG concentration by different physical activities in a patient with type VII glycogenosis. Patient 2 was kept in bed for two days after his admission. On the third day, exercise on a bicycle ergometer was performed before discharge. Blood samples were obtained by venipuncture at the indicated time.](image)

![Fig 3. Effect of ergometer exercise on the concentrations of glycolytic intermediates and adenine nucleotides in erythrocytes from patient 2 with type VII glycogenosis. The blood samples were obtained before (A), immediately after (B), and one hour after (C) exercise. Concentrations are expressed in terms of the ratios to the normal resting values.](image)

![Fig 4. Effect of ammonium and inosine on erythrocyte glycolytic intermediates. Erythrocytes were incubated at a Hct value of 44%. Incubation medium, 100 mmol/L NaCl, 10 mmol/L KCl, 50 mmol/L triethanolamine/HCl, 30 mmol/L glucose, and 1 mmol/L inorganic phosphate (pH 7.4). Concentrations are expressed in terms of the ratios to the controls. (A) O, 25 μmol/L; □, 50 μmol/L; V, 100 μmol/L; □, 150 μmol/L inosine. (B) O, 250 μmol/L; □, 500 μmol/L; □, 1,000 μmol/L ammonium chloride.](image)
similar both in velocity and in kinetics,7 the erythrocytes can be considered to share common metabolic defects. Figure 2 shows that physical activity clearly affected the level of 2,3-DPG in patient 2. During bed rest, 2,3-DPG levels gradually decreased and reached a nadir within one day; however, the increase in 2,3-DPG levels during exercise was very rapid, and the nadir values were similar to those in another family (Table 1).

Analysis of glycolytic intermediates in erythrocytes drawn immediately after exercise showed a marked increase in fructose-1,6-bisphosphate and dihydroxyacetone phosphate plus glyceraldehyde-3-phosphate levels (Fig 3). We also found that a similar increase occurred in types V and III glycogenoses in blood drawn from the antecubital vein after semiischemic forearm exercise (data not shown). Therefore, these alterations in levels of erythrocyte glycolytic intermediates after exercise seem to be common in muscle glycogenoses. These increases in levels of intermediates before and during exercise lasted less than one hour. There were no changes in pH, O2 saturation, or serum inorganic phosphate levels in systemic blood during exercise, which are thought to affect the 2,3-DPG level. It has been reported that alanine is released from muscle but no uptake of branched-chain amino acids occurs in normal subjects during exercise, whereas in type V glycogenosis there is a marked uptake of alanine, leucine, and isoleucine by muscle without a significant release of other amino acids.13,14 In seeking other factors, we demonstrated previously that working muscle released ammonia and inosine to the local circulation in type V and VII glycogenoses4 and also found that even mild exercise could increase plasma ammonia and inosine levels in systemic blood in type VII glycogenosis.15 Ammonia is a potent activator of PFK,7 and inosine freely passes into erythrocytes and is metabolized rapidly through the hexose monophosphate shunt.4 Therefore, we performed an in vitro incubation study to determine the effect of these substances. The addition of ammonium chloride increased the levels of glycolytic intermediates after the PFK step only slightly, but the addition of inosine markedly increased both fructose-1,6-bisphosphate and dihydroxyacetone phosphate plus glyceraldehyde-3-phosphate levels and the 2,3-DPG concentration was also increased. Thus inosine could mainly mimic the effect of exercise in type VII glycogenosis. The concentration of inosine used in vitro may be high (25 to 150 μmol/L), but we observed that inosine in the local blood circulation was increased to more than 20 μmol/L after semiischemic forearm exercise and was metabolized rapidly.15,16 Twenty minutes after incubation with 150 μmol/L inosine, the inosine concentration in the incubation medium decreased to 17.0 μmol/L, although inosine might be produced continuously in vivo unless exercise stopped. Therefore, erythrocytes in vivo could metabolize more inosine than that expected from the in vitro experiment. Although we used normal erythrocytes for the in vitro study, these results lead us to the conclusion that exercise in type VII glycogenosis altered the glycolysis of erythrocytes in circulating blood and that inosine and/or ammonia liberated from muscle is at least partly responsible for this glycolytic alteration. We should take an opportunity to make an erythrocyte incubation study with the patient's erythrocytes.

Inosine and hypoxanthine, which are produced because of excess purine degradation in working muscle and are released into the blood, can be used as precursors for uric acid synthesis in liver. This may contribute to the development of hyperuricemia in types VII, V, and III glycogenoses.12,17,18 The excess purine degradation in muscle during and after mild exercise in type VII glycogenosis19 may result from strenuous exercise in normal humans,20 which is reported also to alter the concentration of erythrocyte 2,3-DPG. These results are controversial; although many investigators21,22 found an increase in 2,3-DPG levels after exercise, others reported a decrease23,24 or an unchanged value.25,26 The many factors involved in strenuous exercise complicate the analysis of erythrocyte glycolysis. Positive factors are considered to be an increased deoxyhemoglobin content,21,22 an increased inorganic phosphate concentration,27 and an increased younger erythrocyte population.22,24 Negative factors are marked increases in lactate levels and acidosis.20,21 These indexes were not altered during this study of type VII glycogenosis.

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