Continuous Renal Replacement Therapy in a 4-year-old Child with Rhabdomyolysis Following Parainfluenza Virus Infection and Hyperammonemia due to Isovaleric Acidemia

Parainfluenza virus infection is one of the causes of fatal rhabdomyolysis. Rhabdomyolysis can be aggravated by mitochondrial fatty acid β-oxidation disorders during prolonged periods of fasting. Moreover, in patients with late-onset isovaleric acidemia, hyperammonemia may occur following catabolic stress. In the present report, we describe a case of a 4-year-old boy with parainfluenza virus infection and late-onset isovaleric acidemia that rapidly progressed to coma, seizures, and cardiorespiratory collapse. His serum ammonia and creatinine kinase (CK) levels were 385 μMol/L and 23,707 IU/L, respectively. Continuous renal replacement therapy (CRRT) was initiated using continuous venovenous hemodiafiltration, after which the ammonia and CK levels returned to normal. Thus, we recommend the immediate initiation of CRRT in the management of patients with life-threatening rhabdomyolysis and hyperammonemia.

Key words: Rhabdomyolysis, Hyperammonemia, Parainfluenza virus, Isovaleric acidemia, Continuous renal replacement therapy

Introduction

Rhabdomyolysis and hyperammonemia due to congenital metabolic disorders are often treated with hemodialysis (HD) [1]. However, conventional HD may be relatively ineffective in removing circulating myoglobin due to its large molecular weight. Continuous renal replacement therapy
CRRT has the advantage of removing relatively large molecules and abolishing rebound hyperkalemia and acidosis through continuous venovenous hemofiltration or hemodiafiltration. Theoretically, convection removes larger molecular-weight solutes better than diffusion [1].

Here, we report a case of rhabdomyolysis and hyperammonemia in a 4-year-old boy following parainfluenza virus infection and late-onset, transient isovaleric acidemia, which was treated successfully with CRRT.

Case report

A 4-year-old boy was transferred to our emergency room due to a change in mental status. Four days before admission, he developed fever, vomiting, and diarrhea. Oral medication was administered at a local clinic, but vomiting and diarrhea continued 5 to 6 times a day. There was no specific perinatal or past medical history. He did not experience any trauma or drug intoxication.

On physical examination, the child was lethargic, stuporous, and severely dehydrated. He had a blood pressure of 100/48 mmHg, a pulse rate of 130/min, a body temperature of 37.5℃, and a respiratory rate of 42/min. His previous body weight of 19 kg had decreased to 18 kg. He did not show any meningeal signs such as Brudzinski or Kernig signs.

Initial laboratory findings revealed hypoglycemia (45 mg/dL, range 70–110), hyperammonemia (385 μMol/L, range 0.0–54), and rhabdomyolysis (creatinine kinase, 1,665 IU/L; range 30–180). Aspartate aminotransferase (AST, 133 IU/L; range 5–40), alanine aminotransferase (ALT, 129 IU/L; range 8–41), lactate dehydrogenase (LDH, 592 IU/L; range 100–200), and uric acid (12.0 mg/dL; range 2.0–5.5) levels were elevated. Prothrombin time was 13.6 seconds (INR 1.18) and activated partial thromboplastin time was 25 seconds. The electrolyte results were as follows: serum sodium of 148 mEq/L (range 135–145), potassium of 3.4 mEq/L (range 3.5–5.5), chloride of 114 mEq/L (range 98–107), and total carbon dioxide of 7 mEq/L (range 24–31), suggesting severe metabolic acidosis. Arterial blood gas analysis results were as follows: pH 7.285, pCO₂ 9.8 mmHg, pO₂ 85.4 mmHg, base excess −18.6 mEq/L, and HCO₃⁻ 4.6 mEq/L. These results also indicated severe metabolic acidosis with respiratory compensation. The calculated anion gap [Na⁺ − (Cl⁻ + HCO₃⁻)] was 27 mEq/L in ambient air. The specific gravity of his urine was elevated to ≥1.035, and the urine was positive for ketone 3. Cerebrospinal fluid profiles were all within normal ranges and no organisms were isolated from blood, urine, or CSF cultures. However, parainfluenza virus was detected by polymerase chain reaction in nasopharyngeal specimens. With a high index of suspicion for organic acidemia, treatment with sodium phenylbutyrate, L-carnitine, lactulose enema, and 10% dextrose in water was begun.

On the third day of admission, laboratory testing revealed rapidly deteriorating rhabdomyolysis. Maximum creatinine kinase (CK) and CK-MB levels were 23,707 IU/L and 131.3 μg/L (range 0.0–5.0), respectively. Tandem mass spectrometry (MS/MS) showed an increase in isovalerylcarnitine [C5] (1.513 μM, range 0.005–0.899), suggesting isovaleric acidemia, which requires a low protein and leucine diet. Increases in AST to 1,626 IU/L, ALT to 691 IU/L, LDH to 2,719 IU/L, uric acid to 10.6 mg/dL (range 0.0–72), and the onset of acute kidney injury (creatinine, 1.0 mg/dL; range 0.3–0.6) indicated CRRT. CRRT was performed for 5 days, after which rhabdomyolysis and hyperammonemia subsided (Fig. 1).

After 2 weeks, urinary aminoacid analysis showed high amounts of β-aminoisobutyric acid (3,103 μmol/g creatinine), arginine (216 μmol/g creatinine), cystathionine (44 μmol/g creatinine), cystine (139 μmol/g creatinine), and ethanolamine (1,837 μmol/g creatinine). Tandem mass spectrometry analysis performed two months later revealed an increase in octanoylcarnitine [C8] (0.803 μM, range 0.010–0.453) and decenoylcarnitine [C10:1] (0.438 μM, range 0.005–0.233), suggesting medium-chain acylcoenzyme dehydrogenase deficiency. However, the A985G and T199C mutations commonly associated with that deficiency had not been detected. He was discharged with puritan (vitamin B2) and L-carnitine syrup. He has been doing well over one year of follow-up and shows no evidence of recurrent rhabdomyolysis or hyperammonemia.
Discussion

Rhabdomyolysis is a syndrome caused by various insults to skeletal muscle and results in the release of myoglobin into the plasma. Massive rhabdomyolysis accompanied by viral myositis and its consequences have been reported by several researchers. In a retrospective series from a pediatric emergency room, 38% of rhabdomyolysis cases were found to have been caused by viral myositis. Several viruses such as adeno, influenza A and B, coxsackie, Epstein–Barr, echo, and measles have been implicated in the pathogenesis of rhabdomyolysis. Parainfluenza virus is known to be one of the causative viruses for complicated rhabdomyolysis. To our knowledge, only 6 cases of rhabdomyolysis induced by parainfluenza virus infection in adolescents and children have been reported in the literature: 3 were attributed to parainfluenza type 1, 2 to parainfluenza type 2, and 1 fatal case was attributed to type 3 (Table 1). While rhabdomyolysis is rare, it is a life-threatening complication of viral myositis. We believe that the most probable cause of rhabdomyolysis in our patient was from a parainfluenza virus infection aggravated by coexistent isovaleric acidemia, which caused severe metabolic acidosis and hyperammonemia, finally resulting in acute kidney injury. CRRT is able to remove myoglobin from the blood in spite of its large molecular weight, thus aborting rebound hyperkalemia and acidosis. Consequently, virus-induced fatal rhabdomyolysis can be effectively treated by CRRT, resulting in successful clinical outcomes.

Isovaleric acidemia, a type of organic acidemia, is an autosomal recessive congenital defect in leucine metabolism caused by deficiency of isovaleryl-CoA dehydrogenase, a mitochondrial enzyme. Most organic acidemias become clinically apparent during shortly after birth or in early infancy. However, there is another phenotype

![Fig. 1](image-url) Changes in serum myoglobin (A) and ammonia (B) levels. Continuous renal replacement therapy effectively decreased both increased serum myoglobin and ammonia levels to normal range.

<table>
<thead>
<tr>
<th>Case</th>
<th>Authors</th>
<th>Age/sex</th>
<th>Virus species</th>
<th>Clinical symptoms</th>
<th>CK</th>
<th>AST</th>
<th>LDH</th>
<th>Serum myoglobin</th>
<th>Serum anion gap</th>
<th>Treatment</th>
<th>Prognosis</th>
</tr>
</thead>
<tbody>
<tr>
<td>Case 1</td>
<td>Pana et al (4) (2011)</td>
<td>10 yr/M</td>
<td>Parainfluenza type 1</td>
<td>Muscle weakness</td>
<td>7,563</td>
<td>71</td>
<td>617</td>
<td>NA</td>
<td>NA</td>
<td>Oral fluid intake</td>
<td>good</td>
</tr>
<tr>
<td>Case 2</td>
<td>Ebbonov et al (6) (2009)</td>
<td>6 yr/F</td>
<td>Parainfluenza type 1</td>
<td>Fever, malaise, sore throat</td>
<td>&gt;50,000</td>
<td>156</td>
<td>NA</td>
<td>9 mEq/L</td>
<td>Hemodialysis</td>
<td>good</td>
<td></td>
</tr>
<tr>
<td>Case 3</td>
<td>Vrsalovic (5) (2007)</td>
<td>5 yr/M</td>
<td>Parainfluenza type 1</td>
<td>Rhinorrhea, vomiting, spastic quadriplegia</td>
<td>22,242</td>
<td>1,040</td>
<td>2,995</td>
<td>NA</td>
<td>NA</td>
<td>CRRT</td>
<td>good</td>
</tr>
<tr>
<td>Case 4</td>
<td>Zvolanek (7) (1984)</td>
<td>8 yr/M</td>
<td>Parainfluenza type 2</td>
<td>Fever, headache, vomiting</td>
<td>4,060</td>
<td>204</td>
<td>NA</td>
<td>NA</td>
<td>NA</td>
<td>Conservative Tx</td>
<td>good</td>
</tr>
<tr>
<td>Case 5</td>
<td>O’Connor et al (8) (1982)</td>
<td>38 yr/M</td>
<td>Parainfluenza type 2</td>
<td>Dark urine</td>
<td>66,000</td>
<td>1,200</td>
<td>4,500</td>
<td>NA</td>
<td>NA</td>
<td>Hydration</td>
<td>good</td>
</tr>
<tr>
<td>Case 6</td>
<td>Ueda et al (9) (1978)</td>
<td>4 yr/M</td>
<td>Parainfluenza type 3</td>
<td>Rhinorrhea, cough</td>
<td>1,700</td>
<td>392</td>
<td>1,032</td>
<td>NA</td>
<td>NA</td>
<td>Aspirin</td>
<td>fatal</td>
</tr>
<tr>
<td>Case 7</td>
<td>Park et al (No type) (2011)</td>
<td>4 yr/M</td>
<td>Fever, vomiting, diarrhea</td>
<td>23,707</td>
<td>1,626</td>
<td>2,719</td>
<td>2,020</td>
<td>27 mEq/L</td>
<td>CRRT</td>
<td>fatal → good</td>
<td></td>
</tr>
</tbody>
</table>

Abbreviations: NA, not available; CK, creatinine kinase; AST, Aspartate aminotransferase; LDH, lactate dehydrogenase
similar to that seen in our case that presents after an initial period of good health in which the patient develops a life-threatening episode of metabolic acidosis characterized by an increased anion gap [12]. This late-onset organic acidemia is difficult to detect. Children with organic acidemia are susceptible to metabolic decompensation during acute episodes of increased catabolism such as intercurrent illness, trauma, surgery, or prolonged episodes of fasting. First-line diagnosis of organic acidemia is by organic acid urine analysis. However, tandem mass spectrometry (MS/MS), which is used in newborn screening, is also helpful in establishing a prompt diagnosis in some unexpected late-onset metabolic crises following catabolic stress [13]. In our patient, catabolic stress due to upper respiratory infection by parainfluenza virus infection, vomiting, diarrhea, and poor oral intake induced the presentation with transient isovaleric acidemia and hyperammonemia and rapidly aggravated the clinical course of rhabdomyolysis. Excessive ammonemia, which occurs in organic acidemia, should be treated early with hemodialysis and medication such as N-carbamylglutamate [10]. CRRT has proved effective in many studies for treating hyperammonemia complicated by congenital metabolic defect [14–16].

In conclusion, this case highlights the significance of maintaining a high index of suspicion for viral infection in children with rhabdomyolysis and emphasizes the role of CRRT in the management of fatal rhabdomyolysis and hyperammonemia in children who have parainfluenza virus infection and coexistent isovaleric acidemia.

Conflict of Interest

The authors declare no potential conflicts of interest with respect to the research, authorship, and/or publication of this article, and no funding.

References


